

Dual Alkali Acceptance Test at Louisville
Gas and Electric Company; Volume II
Appendices D-F

Bechtel Corp.
San Francisco, CA

Prepared for

Industrial Environmental Research Lab.
Research Triangle Park, NC

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DUAL ALKALI ACCEPTANCE TEST
AT LOUISVILLE GAS AND ELECTRIC
COMPANY; VOLUME II.
APPENDICES D-F

Prepared for

Office of Air Quality Planning and Standards
Office of Environmental Engineering and Technology

Prepared by

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Research Triangle Park NC 27711

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16. ABSTRACT The report gives results of the completed acceptance test series run on the dual alkali system serving Louisville Gas and Electric Company's Cane Run Unit 6 boiler. Test results, measuring system performance, included: (1) SO2 removal averaged 94% and 143 ppm outlet concentration; (2) soda ash consumption averaged 0.042 mole soda ash per mole SO2 removed; (3) lime consumption averaged 1.04 mole CaO per mole SO2 removed; (4) power consumption averaged 1.05% of generation; (5) filter cake solids averaged 52.2 wt % insoluble solids; and (6) there was no net particulate matter addition. Various problems attributable to the boiler, the FGD system, and the quality and quantity of the carbide lime supplied to the system delayed acceptance testing until July 1980. The year-long demonstration period officially started in May 1980. The problems experienced and their solutions are discussed. Volume I discusses the test and contains Appendices A-C. This volume (Volume II) contains a discussion of the analytical and sampling methods, quality assurance, and the raw data gathered by Kenvirons. Volume III contains Appendices G-J.			
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Flue Gases	Calcium Oxides	Dual Alkali System	21B
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Tests	Acceptability		14B 14G
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September 1981

DUAL ALKALI ACCEPTANCE TEST
AT LOUISVILLE GAS AND ELECTRIC COMPANY

VOLUME II

Appendices D-F

by

D.A. Watson, T.M. Martin, J.K. Donnelly, N. Zoueshtiagh
Bechtel National, Inc.
50 Beale St.
San Francisco, CA 94119

R.P. Van Ness
Louisville Gas and Electric Company
311 W. Chestnut Street
Louisville, KY 40201

C.R. La Mantia, L.R. Woodland
Arthur D. Little, Inc.
20 Acorn Park
Cambridge, MA 02140

Contract Nos. 68-02-2189 and 68-02-2634

EPA Project Officer: Norman Kaplan

Industrial Environmental Research Laboratory
Office of Environmental Engineering and Technology
Research Triangle Park, N. C. 27711

Prepared For:

U.S. Environmental Protection Agency
Office of Research and Development
Washington, D.C. 20460

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PREFACE

Volumes II and III of this report contain the results of the stack sampling for particulate matter and sulfur dioxide emissions conducted by Kenvirons, Inc. under subcontract to Bechtel National, Inc. Volume II contains Kenviron's test program description, summary of results, field data forms and a portion of the preliminary calculation sheets. Volume III contains the remainder of the calculation sheets and summary forms.

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Appendix D. Kenvirons Tests

1.0 INTRODUCTION

Bechtel National, Inc. has been contracted by the Environmental Protection Agency (Contract No. 68-02-2634) to evaluate the Dual Alkali SO₂ control process on the Louisville Gas and Electric Company Cane Run Station Unit No. 6. Kenvirons, Inc. was subcontracted (Agreement No. 11975-2634-SF-3) to provide testing support services.

The first phase of testing, associated with the demonstration program, consists of an "Acceptance Test". Characteristics of this test, including process monitoring and testing, were presented in a Bechtel report entitled "Acceptance Test Plan", dated April, 1979.

Acceptance testing was originally scheduled for late 1978, then revised to October, 1979. On-going process problems delayed testing on a weekly basis. Testing was scheduled and performed during December 4 through 10, 1979; however, due to unacceptable equipment performance, such testing was designated informational. Kenvirons' test report for that work was submitted to Bechtel on July 9, 1980.

The official acceptance test was thus rescheduled after correcting equipment problems. Testing, which is the basis of this report, was performed during July 17 through 26, 1980.

On the basis of a Bechtel test plan scope, Kenvirons prepared a detailed "Work Plan" dated May, 1980, that described test methods and locations, sampling equipment analysis procedures, etc., in fulfillment of subcontract terms. That plan is incorporated herein by reference since few changes were made.

The Acceptance Test essentially required ten straight days of proper system operation. Financial reimbursement to various participating parties was at stake per the acceptance test. The Kenvirons effort required simultaneous sampling at three different varying locations between the boiler and stack for ten straight days. Particulate and SO₂ mass flows using EPA Method 5-8 were measured at each location. See Table 1 for the test schedule.

Mr. A. Abdulsattar served as Project Manager for Bechtel until June, 1980. Mr. Dave Watson replaced Mr. Abdulsattar as Project Manager prior to testing. Mr. Watson was assisted on-site by Mr. Mickey Martin and Mr. John Donnelly.

Mr. Bob Van Ness served as coordinator for L G & E. Mr. Bud Carter served as project engineer. They were assisted by Mr. Rick Melloan and Ms. Mary Pederson. Mr. Bob Miles and Mr. Terry Rausch served as expeditors. We sincerely appreciate the assistance and hospitality offered by L G & E.

Mr. Norman Kaplan served as Project Officer for EPA.

TABLE 1
TEST SCHEDULE

<u>DAY</u>	<u>DATE</u>	<u>TEAM 1</u>	<u>TEAM 2</u>	<u>TEAM 3</u>
1	7-17-80	STACK	ESP-A Outlet	ESP-B Outlet
2	7-18-80	STACK	ESP-A Outlet	ESP-B Outlet
3	7-19-80	STACK	ESP-A Outlet	ESP-B Outlet
4	7-20-80	STACK	ESP-A Outlet	ESP-B Outlet
5	7-21-80	STACK	ESP-A Outlet	ESP-B Outlet
6	7-22-80	STACK	ESP-A Outlet	ESP-B Outlet
7	7-23-80	STACK	ESP-A Outlet	ESP-B Outlet
8	7-24-80	STACK	ESP-A Outlet	ESP-B Outlet
9	7-25-80	STACK	ESP-A Outlet	ESP-B Outlet
10	7-26-80	STACK	ESP-A Outlet	ESP-B Outlet

Mr. Robert C. Tussey, Jr. served as Principal Investigator for Kenvirons. Dr. Gary Revlett served as analytical coordinator and team leader at the stack. He was assisted by Mr. Parker Moore. Mr. Ron Tussey served as team leader at the ESP-B location. He was assisted by Mr. Paul Ramp and Ms. Donna Blankenship. Mr. Bud Lauritzen served as team leader at the ESP-A location. He was assisted by Mr. Jeff Bralley. Mr. Paul Ramp and Ms. Donna Blankenship, along with Dr. Gary Revlett, performed all sample analysis for Kenvirons. Mr. Bralley performed computer data reduction. Mr. R. S. and R. C. Tussey, Jr. co-authored this report.

2.0 SUMMARY OF RESULTS

Table 2 presents a summary of results for all 30 test runs. As previously indicated, three simultaneous runs were made each day at three different locations by three teams. The following example explains the run code: run BEC-STK-1 is stack run 1 while run BEC-ESP-A-2 is ESP outlet A-side, run number 2 and BEC-ESP-B-3 is ESP outlet B-side, run number 3.

Particulate emission rate is presented as gr/DSCF, gr/ ACF, lb/hour, and lb/MMBTU. SO₂ results are presented as lb/DSCF, lb/ACF, lb/hour, lb/MMBTU and PPM.

Due to intentional variability in process operation each day, we cannot evaluate or discuss in detail, day-to-day comparative test results. Our discussion is limited to probability of accurate and representative samples.

Table 2 presents both sampling data averages and emission calculation results. Particulate emission rates are presented as grains per dry standard cubic foot (Gr/DSCF), pounds per DSCF, grains per actual cubic foot (Gr/ACF), pounds per hour (LB/Hr), and pounds per million British Thermal Units (LB/MMBTU). Sulfur dioxide emission rates are presented as (LB/MMBTU), and parts per million standard dry (PPM) and parts per million stack conditions.

Table 3 presents a summary (from Table 2) of results, average and range, for pertinent variables. Most results were within expected ranges. A brief discussion of averages and ranges follows.

*Our isokinetic sampling rate was within the method-specified range of 100% $\pm$ 10% for all runs except STK002 (11.6%). The slightly out of range rate that day was due to a significantly higher % Moisture content (15.0%) compared to the first run at the stack (8.8%). Since the rate was only 1.6% out of range, the results should not be significantly affected.

*The moisture content at the stack was low for day 1 (8.8%) but higher and consistent for the remaining days (13.7% - 15%). ESP A and B side moisture average was within about one-half percent and the range was less than three percent.

*Oxygen content at the stack was consistent (7.7% - 8.8%) as measured by Orsat. ESP A and B values were consistent, varying from 6.8% to 9.3% using a Fyrite. ESP A and B averages were very close at 7.75% and 7.88% respectively. The slightly higher stack value indicates air inleakage.

*Temperature at the stack varied from 167.8°F to 179.2°F, which is a reasonably low range. The ESP A average was higher (304.4°F) than the ESP B value (293.47°F) by about 12°F. The ESP B range was 25°F and the ESP A range was 13°F.

*ESP B velocity (2099 FPM) was lower than ESP A (2232 FPM). ESP B range was higher (500) than ESP A (200 FPM). Stack velocity range was 300 FPM.

OUTPUT DATE - 08/27/80
TIME - 14130151

BECHTEL 1980 DEMO-ACCEPTANCE TEST - LOISE CANE RUN NO.4

TABLE 2
TEST RESULTS SUMMARY

KENVIRONS, INC.
PROJECT NO. B0019

PARTICULATE AND SO2 EMISSION DATA

NAME	DESCRIPTION	UNITS	1	2	3
IN	DATE OF RUN		ESPA01	STK001	ESPA02
IN	TIME OF RUN		07/17/80	07/17/80	07/18/80
IN	PROBE TIP DIAMETER	IN	1428-1724	1428-1655	1100-1338
IN	NET TIME OF RUN	MIN	0.238	0.241	0.238
IN	NET TIME OF RUN	MIN	100.0	100.0	100.0
IN	NUMBER OF SAMPLING POINTS		50	50	50
IN	DRY GAS METER CALIB. FACTOR		1.0240	1.0370	1.0250
IN	BAROMETRIC PRES, LAB	IN.HG	29.66	29.66	29.80
IN	BAROMETRIC PRES, CONSOLE	IN.HG	29.61	29.61	29.75
IN	AUG DRY PRES DROP	IN.H2O	0.580	0.706	0.640
IN	VOL DRY GAS-METER COND	DECF	42.60	48.70	44.11
IN	AUG GAS METER TEMP	DEG.F	104.8	124.4	104.1
IN	VOL DRY GAS-STD COND	SCF	40.41	45.07	42.14
IN	TOTAL H2O COLLECTED	ML	81.1	89.5	91.4
IN	VOL H2O VAPOR-STD COND	SCF	3.82	4.21	4.31
IN	PERCENT MOISTURE BY VOLUME		8.4	8.5	9.3
IN	HOLE FRACTION DRY GAS		0.914	0.915	0.907
IN	PERCENT CO2 BY VOLUME, DRY		12.0	12.0	12.0
IN	PERCENT O2 BY VOLUME, DRY		7.7	7.3	7.5
IN	PERCENT CO BY VOLUME, DRY		0.0	0.0	0.0
IN	PERCENT N2 BY VOLUME, DRY		80.3	80.7	80.5
IN	MOLECULAR WT-DRY STK GAS		30.23	30.21	30.22
IN	MOLECULAR WT-DRY STK GAS		29.17	29.17	29.09
IN	PIDOT TUBE COEFFICIENT		0.840	0.840	0.840
IN	AUG SORT STK VEL HEAD	IN.H2O	0.522	0.560	0.548
IN	AVG STACK TEMPERATURE	DEG.F	300.1	283.4	311.5
IN	STACK PRESSURE, ABSOLUTE	IN.HG	28.46	28.50	28.44
IN	AVG STACK GAS VELOCITY	FPM	2150	2280	2270
IN	STACK AREA	IN2	33618	33618	33618
IN	STK FLOWRATE, DRY, STD	SCFM	303000	327000	315000
IN	ACTUAL STACK FLOWRATE	ACFM	502000	532000	530000
IN	PERCENT ISOINETIC		100.7	100.9	101.0
IN	PARTICULATE WT-TOTAL	LB	628.70	942.90	822.10
IN	PART. LOAD-TTL, STD	LB/SCF	0.240	0.330	0.301
IN	PART. LOAD-TTL, STD	LB/SCF	0.145	0.204	0.179
IN	PART. LOAD-TTL, STD	LB/SCF	0.145	0.204	0.179
IN	PART. LOAD-TTL, STD	LB/SCF	0.145	0.204	0.179
IN	EMISSION RATE	LB/HRT	624.0000	930.0000	813.0000
IN	F FACTOR		0.5320	0.7120	0.6590
IN	SCONCA SO2, CONC., STD	PPM	3.84E-04	3.84E-04	3.57E-04
IN	SCONCA SO2, CONC., STD	PPM	2.33E-04	2.39E-04	2.12E-04
IN	SCONCA SO2 EMISSION RATE	LB/HRT	7028.138	7627.597	6752.409
IN	SCONCA SO2 EMISSION RATE	LB/HRT	5.990	5.840	5.470
IN	SCONCA SO2, CONC., STD	PPM	2330	2330	2150
IN	SCONCA SO2, CONC., STD	PPM	2130	2130	1950
IN	STK001		0.0000527	0.0000527	0.0000480
IN	STK002		0.0000480	0.0000480	0.0000480
IN	STK003		0.0000480	0.0000480	0.0000480
IN	STK004		0.0000480	0.0000480	0.0000480
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IN	STK100		0.0000480	0.0000480	0.0000480
IN	STK101		0.0000480	0.0000480	0.0000480
IN	STK102		0.0000480	0.0000480	0.0000480
IN	STK103		0.0000480	0.0000480	0.0000480
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IN	STK108		0.0000480	0.0000480	0.0000480
IN	STK109		0.0000480	0.0000480	0.0000480
IN	STK110		0.0000480	0.0000480	0.0000480
IN	STK111		0.0000480	0.0000480	0.0000480
IN	STK112		0.0000480	0.0000480	0.0000480
IN	STK113		0.0000480	0.0000480	0.0000480
IN	STK114		0.0000480	0.0000480	0.0000480
IN	STK115		0.0000480	0.0000480	0.0000480
IN	STK116		0.0000480	0.0000480	0.0000480
IN	STK117		0.0000480	0.0000480	0.0000480
IN	STK118		0.0000480	0.0000480	0.0000480
IN	STK119		0.0000480	0.0000480	0.0000480
IN	STK120		0.0000480	0.0000480	0.0000480
IN	STK121		0.0000480	0.0000480	0.0000480
IN	STK122		0.0000480	0.0000480	0.0000480
IN	STK123		0.0000480	0.0000480	0.0000480
IN	STK124		0.0000480	0.0000480	0.0000480
IN	STK125		0.0000480	0.0000480	0.0000480
IN	STK126		0.0000480	0.0000480	0.0000480
IN	STK127		0.0000480	0.0000480	0.0000480
IN	STK128		0.0000480	0.0000480	0.0000480
IN	STK129		0.0000480	0.0000480	0.0000480
IN	STK130		0.0000480	0.0000480	0.0000480
IN	STK131		0.0000480	0.0000480	0.0000480
IN	STK132		0.0000480	0.0000480	0.0000480
IN	STK133		0.0000480	0.0000480	0.0000480
IN	STK134		0.0000480	0.0000480	0.0000480
IN	STK135		0.0000480	0.0000480	0.0000480
IN	STK136		0.0000480	0.0000480	0.0000480
IN	STK137		0.0000480	0.0000480	0.0000480

TABLE 2 (Cont.)

KENVIRONS, INC.
PROJECT NO. 80019OUTPUT DATE - 08/27/80
TIME - 14130151

BECHTEL 1980 DEMO-ACCEPTANCE TEST - LGIE CANE RUN NO.6

PARTICULATE AND SO2 EMISSION DATA

NAME	DESCRIPTION	UNITS	ESPA04	ESP004	STK004	ESPA05	ESP005	STK005	ESPA06	ESP006	STK006
DM	TITLE OF RUN		07/20/80	07/20/80	07/20/80	07/21/80	07/21/80	07/21/80	07/22/80	07/22/80	07/22/80
TT	DATE OF RUN		1230-1454	1230-1448	1235-1424	1030-1243	1030-1233	1030-1219	1104-1338	1109-1416	1035-1216
NP	PROBE TIP DIAMETER	IN	0.238	0.241	0.172	0.238	0.241	0.172	0.238	0.241	0.172
Y	NET TIME OF RUN	MIN	100.0	100.0	84.0	100.0	100.0	84.0	100.0	100.0	84.0
Y	NUMBER OF SAMPLING POINTS		50	50	28	50	50	28	50	50	28
Y	DRY GAS METER CALIB. FACTOR		1.0245	1.0355	1.0155	1.0250	1.0350	1.0155	1.0245	1.0350	1.0155
Y	BAROMETRIC PRES. LAB	IN.HG	29.85	29.85	29.85	29.88	29.88	29.88	29.79	29.79	29.79
Y	BAROMETRIC PRES. CONSOLE	IN.HG	29.80	29.80	29.83	29.83	29.83	29.86	29.74	29.74	29.77
Y	AVG DRY PRES DROP	IN.H2O	0.651	0.634	0.478	0.648	0.613	0.482	0.607	0.613	0.440
Y	VOL DRY GAS-METER COND	DCF	44.76	45.23	34.49	45.96	44.19	33.86	44.24	43.12	32.82
Y	VOL DRY GAS-METER TEMP	DEG.F	118.1	121.7	118.6	110.9	116.5	115.2	89.4	98.7	103.2
Y	VOL DRY GAS-STD COND	DSCF	41.77	42.40	31.69	43.50	41.82	31.32	43.36	41.98	30.88
Y	TOTAL H2O COLLECTED	NL	94.0	68.5	115.7	102.5	97.3	115.1	108.6	101.8	111.7
Y	VOL H2O VAPOR-STD COND	SCF	4.52	3.22	5.45	4.82	4.58	5.42	5.11	4.79	5.26
Y	PERCENT MOISTURE BY VOLUME		9.8	7.1	14.7	10.0	9.9	14.7	10.5	10.2	14.5
Y	MOLE FRACTION DRY GAS		0.902	0.927	0.853	0.900	0.901	0.853	0.895	0.898	0.855
Y	PERCENT CO2 BY VOLUME, DRY		12.8	12.1	10.8	12.8	13.0	10.7	11.1	11.5	11.1
Y	PERCENT CO BY VOLUME, DRY		8.5	9.0	8.2	7.8	8.3	8.1	8.1	8.0	7.7
Y	PERCENT N2 BY VOLUME, DRY		0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Y	MOLECULAR WT-DRY STK GAS		28.9	28.9	81.0	29.5	28.7	81.2	30.1	30.1	81.2
Y	MOLECULAR WT-DRY STK GAS		30.38	30.30	30.04	30.35	30.41	30.03	30.10	30.16	30.08
Y	PITOT TUBE COEFFICIENT		29.17	29.33	28.29	29.12	29.19	28.26	28.83	28.92	28.32
Y	AVG SORT STK VEL HEAD	IN.H2O	0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840
Y	AVG STACK TEMPERATURE	DEG.F	0.541	0.504	0.855	0.543	0.500	0.855	0.546	0.546	0.523
Y	STACK PRESSURE, ABSOLUTE	IN.HG	301.2	280.4	179.2	308.9	305.3	174.3	298.0	300.8	175.9
Y	AVG STACK GAS VELOCITY	FT/M	28.75	28.75	29.57	28.73	28.73	29.60	28.67	28.67	29.51
Y	STACK AREA	IN2	2250	2030	3320	2320	2040	3210	2250	2160	3200
Y	STK FLOWRATE, DRY, STD	DSCFM	33618	33618	40828	33618	33618	40828	33618	33618	40828
Y	ACTUAL STACK FLOWRATE	ACFM	311000	302000	436000	322000	287000	438000	314000	301000	435000
Y	PERCENT ISOKINETIC		518000	474000	913000	542000	480000	909000	526000	504000	907000
Y	PARTICULATE WT-TOTAL	LB	1080.70	899.40	91.20	1397.10	1068.40	90.30	1212.50	1788.50	91.30
Y	PART. LOAD-TTL, STD	LB/DSCF	0.399	0.327	0.044	0.496	0.394	0.044	0.431	0.457	0.046
Y	PART. LOAD-TTL, STD	LB/ACF	0.240	0.208	0.031	0.294	0.235	0.031	0.258	0.392	0.032
Y	PART. EMIS-TOTAL	LB/HR	1066.0345	847.0000	242.0000	1367.5283	969.0000	243.0000	1161.2044	1695.5126	248.0000
Y	EMISSION RATE	LB/HMSTU	0.9420	0.8040	0.1030	1.1100	0.9200	0.1030	0.9900	1.4900	0.1030
Y	F FACTOR	DSCF/HMSTU	9820	9820	9820	9820	9820	9820	9820	9820	9820
Y	SO2, CONC., STD	LB/DSCF	3.84E-04	4.10E-04	2.43E-05	3.87E-04	3.99E-04	2.27E-05	3.90E-04	4.11E-04	4.11E-05
Y	SO2, CONC., STD	LB/ACF	2.38E-04	2.81E-04	1.83E-05	2.30E-04	2.38E-04	1.59E-05	2.33E-04	2.45E-04	2.88E-05
Y	SO2 EMISSION RATE	LB/HR	7394.102	7437.808	1003.707	7471.553	6860.159	870.000	7343.135	7420.226	1544.731
Y	SO2 EMISSION RATE	LB/HMSTU	6.550	7.080	0.425	6.040	6.510	0.364	6.260	6.540	0.441
Y	SO2, CONC., STD	PPM	2390	2480	159	2330	2410	137	2350	2480	248
Y	SO2, CONC., STD	PPM	2150	2300	135	2100	2170	117	2100	2230	212

OUTPUT DATE - 08/27/80
TIME - 1411:45

1980 DEMO-ACCEPTANCE TEST

PARTICULATE AND SO₂ EMISSION DATA

NAME	DESCRIPTION	UNITS	ESPA07	ESPR07	STK007	ESFA08	ESP808	STK008	ESPA09	ESPR09	STK009
	TITLE OF RUN		07/23/80	07/23/80	07/23/80	07/24/80	07/24/80	07/24/80	07/25/80	07/25/80	07/25/80
	DATE OF RUN		1620-1832	1620-1923	1620-1814	1100-1310	1100-1302	1100-1244	1145-1334	1145-1330	1145-1331
	TIME OF RUN		0.238	0.241	0.172	0.238	0.241	0.172	0.238	0.241	0.172
DN	PROBE TIP DIAMETER	IN	100.0	100.0	84.0	100.0	100.0	84.0	100.0	100.0	84.0
TY	WET TIME OF RUN	MIN	50	50	28	50	50	28	50	50	28
NY	NUMBER OF SAMPLING POINTS		1.0230	1.0320	1.0115	1.0245	1.0340	1.0130	1.0240	1.0338	1.0135
YP	DRY GAS METER CALIB. FACTOR		29.80	29.80	29.80	29.82	29.82	29.82	29.84	29.84	29.84
FP	BAROMETRIC PRES., LAB	IN.HG	29.75	29.75	29.58	29.77	29.77	29.60	29.79	29.79	29.79
PBC	BAROMETRIC PRES., CONSOLE	IN.HG	0.535	0.450	0.364	0.407	0.363	0.397	0.590	0.542	0.437
PM	AUG DRF PRES DROP	IN.H2O	41.87	39.70	29.43	44.31	42.69	30.98	43.53	42.35	33.01
PM	AUG DRY GAS-METER COND	DCF	94.7	111.8	102.2	93.5	107.9	100.3	98.9	113.4	104.7
TH	AUG GAS METER TEMP	DEG.F	40.58	37.85	27.66	43.14	40.89	29.28	41.99	40.17	31.09
VMST	VOL DRY GAS-STD COND	SCF	90.0	80.8	93.2	107.5	99.3	109.1	95.9	87.2	110.0
VM	TOTAL H2O COLLECTED	ML	4.24	3.80	4.39	5.06	4.67	5.14	4.51	4.10	5.18
VWV	VOL H2O VAPOR-STD COND	SCF	9.5	9.2	13.7	10.5	10.3	15.9	9.7	9.3	14.3
VWV	PERCENT MOISTURE BY VOLUME		0.903	0.908	0.863	0.895	0.851	0.897	0.903	0.907	0.857
PHOS	PERCENT CO2 BY VOLUME, DRY		10.7	11.5	10.5	11.2	11.5	10.8	11.1	10.7	7.9
MD	MOLE FRACTION DRY GAS		7.3	7.8	7.8	7.3	7.2	7.7	8.5	9.3	7.9
PC02	PERCENT CO2 BY VOLUME, DRY		0.0	0.0	0.1	0.0	0.0	0.0	0.0	0.0	0.0
PC02	PERCENT O2 BY VOLUME, DRY		82.0	80.7	81.5	81.3	81.4	81.4	80.4	81.5	81.4
PC0	PERCENT N2 BY VOLUME, DRY		30.00	30.15	30.00	30.08	30.13	30.04	30.12	29.85	30.03
PH2	PERCENT H2 BY VOLUME, DRY		28.87	29.04	28.36	28.81	28.88	28.25	28.75	28.31	28.75
MVD	MOLECULAR WT-DRY STK GAS		0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840
MVD	MOLECULAR WT-STK GAS		0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840	0.840
MU	PITOT TUBE COEFFICIENT		0.514	0.449	0.783	0.549	0.512	0.821	0.541	0.472	0.860
CP	AUG SORT STK VEL HEAD	IN.H2O	304.5	284.6	170.7	309.3	294.9	167.8	300.4	294.8	174.7
DP8	AUG STACK TEMPERATURE	DEG.F	28.80	28.82	29.52	28.72	28.65	29.35	28.74	28.74	29.57
TS	STACK PRESSURE, ABSOLUTE	IN.HG	2130	2930	2930	2280	2110	3070	2230	3220	3220
PS	AUG STACK GAS VELOCITY	FPM	33618	33618	40828	33618	33618	40828	33618	33618	40828
US	STACK AREA	IN2	300000	264000	593000	314000	294000	614000	313000	288000	444000
AS	STK FLOWRATE, DRY, STD	SCFM	498000	426000	832000	537000	492000	849000	521000	473000	914000
GS	ACTUAL STK FLOWRATE	ACFM	102.4	105.0	97.6	103.9	102.0	99.7	101.2	102.7	100.7
GA	PARTICULATE ISOKINETIC		718.50	921.70	81.10	808.50	805.40	76.30	907.40	1242.00	98.40
PERI	PARTICULATE WT-TOTAL	LB/SCF	0.000033900	0.000050646	0.00000646	0.00004130	0.00004340	0.00000574	0.00004740	0.00004820	0.00000699
MT	PART. LOAD-TTL, STD	LB/SCF	0.144	0.234	0.032	0.170	0.183	0.028	0.201	0.291	0.035
CAN	PART. LOAD-TTL, STD	SCFM	701.0000	854.0000	230.0000	778.0000	770.0000	212.0000	876.0000	1178.8716	270.0000
C	PART. LOAD-TTL, STD	LB/ACF	0.5896	0.8470	0.1020	0.4250	0.4490	0.0893	0.7890	1.2000	0.1100
CAT	PART. LOAD-TTL, STD	LB/HRTU	9820	9820	9820	9820	9820	9820	9820	9820	9820
E	EMISSION RATE	LB/HRTU	3.39E-04	3.37E-04	2.27E-03	3.52E-04	3.47E-04	2.15E-05	4.19E-04	4.04E-04	2.21E-03
F	F FACTOR		2.03E-04	2.09E-04	1.62E-05	2.08E-04	2.09E-04	1.52E-05	2.52E-04	2.47E-04	1.54E-03
SCONC	S02 CONC., STD	LB/SCF	6073.866	5338.111	807.000	6629.854	6158.371	794.000	7876.590	7019.456	854.000
SCONCA	S02 CONC., STK	LB/HRTU	5.100	5.250	0.357	5.330	5.190	0.335	6.930	7.150	0.349
SCHR	S02 EMISSION RATE	SCFM	2040	2700	137	2120	2100	130	2530	2450	131
SCHM	S02 EMISSION RATE	PPH	1850	1840	118	1900	1880	111	2280	2220	111
SCPPH	S02 CONC., STD										
SCPPHA	S02 CONC., STK										

TABLE 2 (Cont.)

OUTPUT DATE - 08/27/80
 TIME - 14:33:45
 KENWIRONS, INC.
 PROJECT NO. 80019
 BECHTEL 1980 DEMO-ACCEPTANCE TEST - LOBE CANE RUN NO. 6

PARTICULATE AND SO2 EMISSION DATA

NAME		DESCRIPTION	UNITS	ESP810		ESP810	STK810
		TITLE OF RUN		07/26/80	07/26/80	07/26/80	07/26/80
		DATE OF RUN		0930-1138	0930-1129	0935-1114	
DN	TT	PROBE TIP DIAMETER	IN	0.238	0.241	0.172	
NP	Y	NET TIME OF RUN	MIN	100.0	100.0	84.0	
		NUMBER OF SAMPLING POINTS		50	50	28	
PR		DRY GAS METER CALIB. FACTOR		1.0235	1.0340	1.0130	
PRC		BAROMETRIC PRES., LAB	IN.HG	29.78	29.78	29.78	
PH		BAROMETRIC PRES., CONSOLE	IN.HG	29.73	29.73	29.56	
PH		AVG DRF PRES DROP	IN.H2O	0.561	0.550	0.398	
TM		VOL DRY GAS-METER COND	SCF	42.84	41.53	31.38	
TM		AVG GAS METER TEMP	DEG.F	99.4	108.3	105.1	
VMSTD		VOL DRY GAS-STD COND	SCF	41.18	39.70	29.37	
VW		TOTAL H2O COLLECTED	ML	92.9	84.5	100.6	
VWV		VOL H2O VAPOR-STD COND	SCF	4.37	4.07	4.74	
PHOS		PERCENT MOISTURE BY VOLUME		9.4	9.3	13.9	
ND		HOLE FRACTION DRY GAS		0.904	0.907	0.861	
PCO2		PERCENT CO2 BY VOLUME, DRY		11.5	12.8	10.7	
PCO		PERCENT CO BY VOLUME, DRY		7.0	7.3	7.9	
PN2		PERCENT N2 BY VOLUME, DRY		0.0	0.0	0.1	
MUD		MOLECULAR WT-DRY STK GAS		81.5	80.0	81.3	
MU		MOLECULAR WT-STK GAS		30.12	30.33	30.03	
CP		PITOT TUBE COEFFICIENT		28.94	29.18	28.36	
DPS		AVG SORT STK VEL HEAD	IN.H2O	0.840	0.840	0.840	
TS		AVG STACK TEMPERATURE	DEG.F	0.523	0.499	0.823	
PS		STACK PRESSURE, ABSOLUTE	IN.HG	300.4	291.9	172.5	
VS		AVG STACK GAS VELOCITY	FT/HR	28.57	28.44	29.50	
AS		STACK AREA	IN2	2160	2040	3080	
OS		STK FLOWRATE, DRY, STD	ACFM	33618	33618	40828	
GA		ACTUAL STACK FLOWRATE	ACFM	305000	290000	420000	
MT		PERCENT ISOKEINETIC		505000	476000	874000	
CAN		PARTICULATE WT-TOTAL	LB	102.9	100.9	99.2	
C		PART. LOAD-TTL, STD	GR/DSCF	1216.76	710.20	86.00	
CAT		PART. LOAD-TTL, STD	LB/DSCF	0.456	0.274	0.045	
CAU		PART. LOAD-TTL, STD	GR/ACF	.00006310	.00003940	.00000445	
E		EMISSION RATE	LB/HR	0.273	0.168	0.032	
F		F FACTOR	LB/HMBTU	1181.9492	484.0000	240.0000	
SCONC		SO2, CONC., STD	DSCF/HMBTU	0.920	0.5730	0.1020	
SCONCA		SO2, CONC., STK	LB/DSCF	9820	9820	9820	
SCHR		SO2 EMISSION RATE	LB/ACF	4.00E-04	3.92E-04	2.27E-03	
SCHH		SO2 EMISSION RATE	LB/HR	2.40E-04	2.19E-04	1.61E-03	
SCPPM		SO2, CONC., STD	PPM	7256.073	6816.098	845.000	
SCPPMA		SO2, CONC., STK	PPM	5.900	5.890	0.359	
				2410	2360	137	
				2180	2140	118	

TABLE 3
RESULTS AVERAGE AND RANGE

DESCRIPTION	AVERAGE			RANGE		
	ESP-A	ESP-B	STACK	ESP-A	ESP-B	STACK
1. % Moisture	9.71	9.16	13.86	8.60 - 10.50	7.10 - 10.30	8.80 - 15.00
2. % O ₂	7.75	7.88	8.11	7.00 - 8.50	6.80 - 9.30	7.70 - 8.80
3. Stack Temperature (°F)	304.40	293.47	173.51	298.0 - 311.5	280.4 - 305.3	167.8 - 179.2
4. Velocity (FPM)	2232	2099	3159	2130 - 2320	1820 - 2340	2930 - 3330
5. Flow Rate (DSCFM)	311,500	298,600	634,600	300,000 - 322,000	264,000 - 335,000	593,000 - 666,000
6. Flow Rate (ACFM)	521,300	489,200	895,800	498,000 - 542,000	426,000 - 545,000	832,000 - 945,000
7. % Isokinetic	102.2	103.9	102.8	100.7 - 104.3	100.9 - 107.5	97.6 - 111.6
8. Particulate						
GR/DSCF	0.364	0.380	0.044	0.240 - 0.496	0.174 - 0.657	0.037 - 0.049
LB/HR	975.58	975.39	237.50	624 - 1367	438 - 1695	211 - 270
LB/MMBTU	0.8147	0.8642	0.1004	0.5320 - 1.110	0.3620 - 1.490	0.0893 - 0.1110
9. SO ₂						
LB/HR	7088	6868	961	6073 - 7876	5338 - 7645	794 - 1565
LB/MMBTU	5.93	6.07	0.405	5.10 - 6.93	5.19 - 7.15	0.335 - 0.641
PPM (Dry Std.)	2266	2311	152	2040 - 2530	2030 - 2480	124 - 248

*Dry standard flow rate at the ESP B was less (298,600) than at the ESP A (311,500). Less total flow ($311,500 + 298,600 = 610,100$ DSCFM) was measured at the ESP than stack (634,600). The difference was less than 4% indicating good sampling and measuring performance.

*The average particulate emission rate at the ESP A and B was very close at 0.815 and 0.864 LB/MMBTU. The range at both locations, however, was very high (0.5 at A and 1.1 at B). The stack average was 0.1004 LB/MMBTU, which is essentially at the 1971 NSPS. The stack range was reasonably low (0.089 to 0.111). The average scrubber particulate collection efficiency was 88%. ESP outlet values are much higher than expected.

*The average SO_2 emission rate at the ESP A and B was very close at 5.93 and 6.07 LB/MMBTU. The stack rate was very low, averaging 0.405 LB/MMBTU, which is significantly below the 1971 NSPS (1.2). This gives an approximate scrubber SO_2 efficiency of 93%.

All calculations in Table 2 were performed once manually; then all data were entered into computer for final confirming calculation. Differences were resolved such that the final computer results should be essentially correct.

Appendix A presents copies of all raw field data forms; Appendix B presents particulate analytical results; Appendix C presents SO_2 analytical data; Appendix D presents particulate calculations; Appendix E presents computer calculations and Appendix F presents equipment calibrations.

3.0 TESTING PROCEDURES

3.1 Sample Locations

All testing was conducted at three locations in the ductwork between the ESP and stack (see Table 1). Figure 1 presents a plan view of the ESP, scrubber, and stack. The gas stream is split into two equal parts as it leaves the boiler. Thus, the ESP and scrubber consist of an "A" side (north) and "B" side (south) as illustrated. The ducts are connected to a common plenum after the ESP (before scrubber) to allow pressure balancing. Also, this plenum serves as a scrubber bypass duct. The three test locations are:

1. ESP Outlet "A"
2. ESP Outlet "B"
3. Stack

EPA Method 1 was used as a general guide in selecting the number and location of sample points. At the ESP location, compromises were made due to the lack of straight ductwork prior to ports. Generally, we targeted a minimum of 2 minutes per point, and approximately 50 points per duct for non Method 1 locations. Also, the sample representative area per point (length and width) dimension ratio should not exceed 2:1. The sample time per point was adjusted to allow a total time at all three locations for the day to be approximately simultaneous. Table 4 presents a summary of test location characteristics.

TABLE 4

<u>Sample Location</u>	<u>No. Ports</u>	<u>Points Per Ports</u>	<u>Total Points</u>	<u>Minutes Per Point</u>	<u>Total Time (minutes)</u>
ESP Outlet A	10	5	50	2	100
ESP Outlet B	10	5	50	2	100
Stack	4	7	28	3	84

3.1.1 ESP Outlet (A and B)

The ESP outlet duct "A" is identical to duct "B". Figure 2 illustrates a side view of the ESP outlet. Three straightening vanes turn the gases from a horizontal flow to a down flow direction past the test ports. The 10-foot long sample probe entered horizontally. Figure 3 illustrates a face view of the duct with 10 ports (4-inch pipe with caps) on each of the A and B sides. A straightening vane stops a few inches prior to the port elevation and three inches horizontally. Dust blowoff from the vane could cause sample contamination; however, we had no expedient method of evaluating the problem. Figure 2 also illustrates the distance to all five sample points. The building roof provided a convenient work access area for all ports.

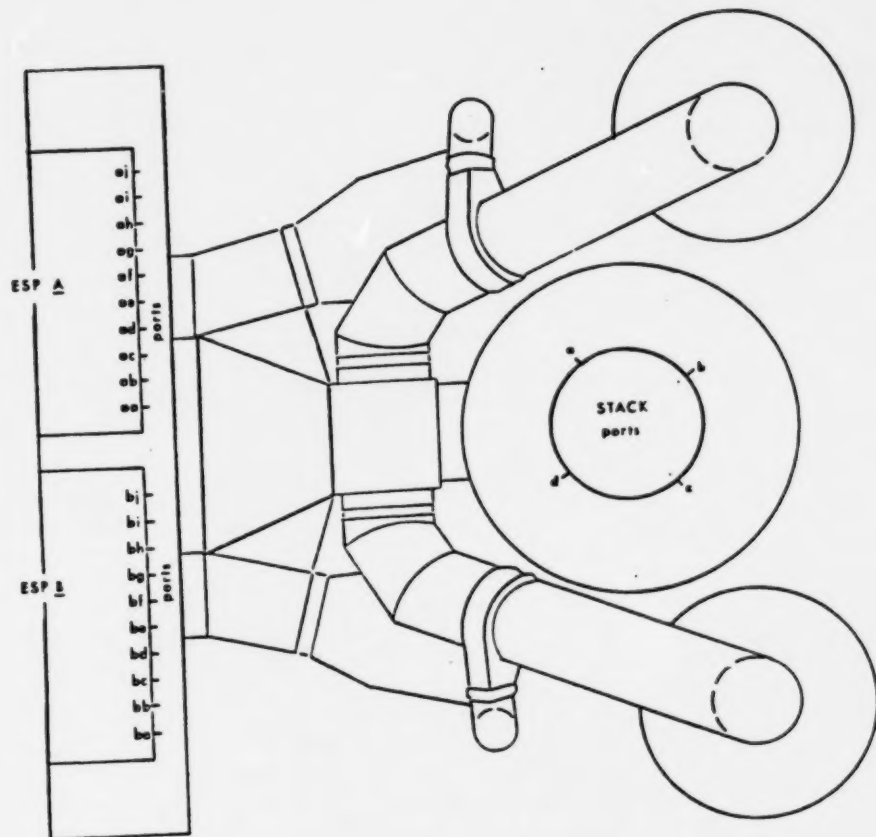


FIGURE 1

PLAN VIEW OF TEST LOCATIONS

3.1.2 Stack

Characteristics of the test location at the stack are presented in Figures 4 and 5. Figure 4 presents a side view and Figure 5 shows a cross-section. The port elevation is 715 feet and the breeching top is at approximately 493 feet. Considering a diameter of 19 feet, we thus have approximately 11-plus diameters to the test ports. EPA Method 1 would thus allow 12 total sample points based on an upstream disturbance (distance B in Method 1). The top of the stack is at elevation 962 feet (approximately), which provides significantly more than 2 diameters (distance A in Method 1); thus, 12 points would be acceptable. For improved accuracy we used 28 points or 7 each for the 4 ports.

Access to the sample platform was gained by climbing an offset ladder approximately 250 feet. The actual sampling process was tedious since the openings in the concrete shell were just barely large enough to pass the test equipment. L G & E provided an elevator between the liner and shell to facilitate equipment handling.

3.2 Sampling and Analysis Methods

3.2.1 Particulate and SO₂ Sampling

A combination EPA Method 5-8 sample train was used to measure particulates and SO₂ simultaneously (Federal Register, August 18, 1977). The particulate sample was isokinetically obtained per Method 5. The standard method 5 condenser setup was modified and a method 8 impinger train substituted, including a filter between the first and second impinger. The H₂O₂ concentration was varied in the second and third impingers depending on expected SO₂ concentration. 100 ml of 3% H₂O₂ concentration was used at the stack and 100 ml of 12% H₂O₂ was used at the ESP-A and B. Approximately 200 grams of silica gel was used in the last (4th) impinger. After the solutions were measured by graduated cylinder and poured into the impingers the solution plus impinger was weighed to the nearest 0.1 gram using an Ohaus 310 double beam balance.

The sampling trains used were Lear-Siegler, Inc. Model PM/100 Manual Stack Samplers (see Figure 6). The sampling trains consisted of a buttonhook-type nozzle, the appropriate probe, a two-section sample box, a 50' or 75' umbilical cord, and a control console. The control console contained a leak-free diaphragm pump, a dry gas meter, coarse and fine adjust valves, an inclined manometer, two thermometers, a pyrometer, a vacuum gauge, and umbilical cord connections. The diaphragm pump moved the stack gases through the sampling system while the dry gas meter measured the volume of gas sampled. Gas flow through the system was controlled with the coarse and fine adjust valves. The range of the dual inclined manometer was 0-5 in. H₂O graduated in 0.01 in. H₂O divisions from 0-1" and 0.1 in. H₂O from 1"-5". The manometer provided for simultaneous measurement of the pressure drop across the orifice in the control console and the differential pressure of the stack gas velocity at the sampling point(s). The umbilical cord and connectors united the metering/control console with the sample box. One portion of the sample box held a heating element which maintained the temperature around the filter holder at approximately 250°F. The other side of

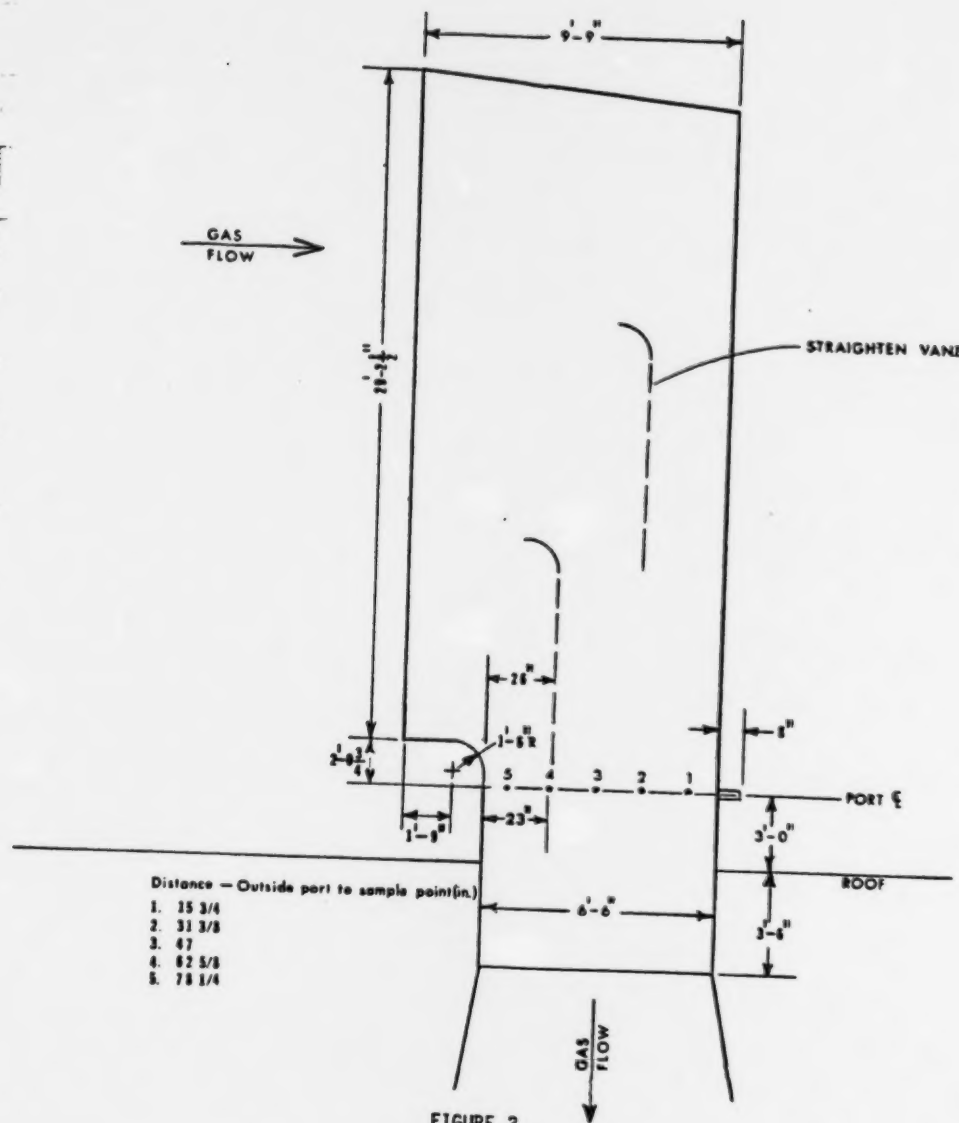
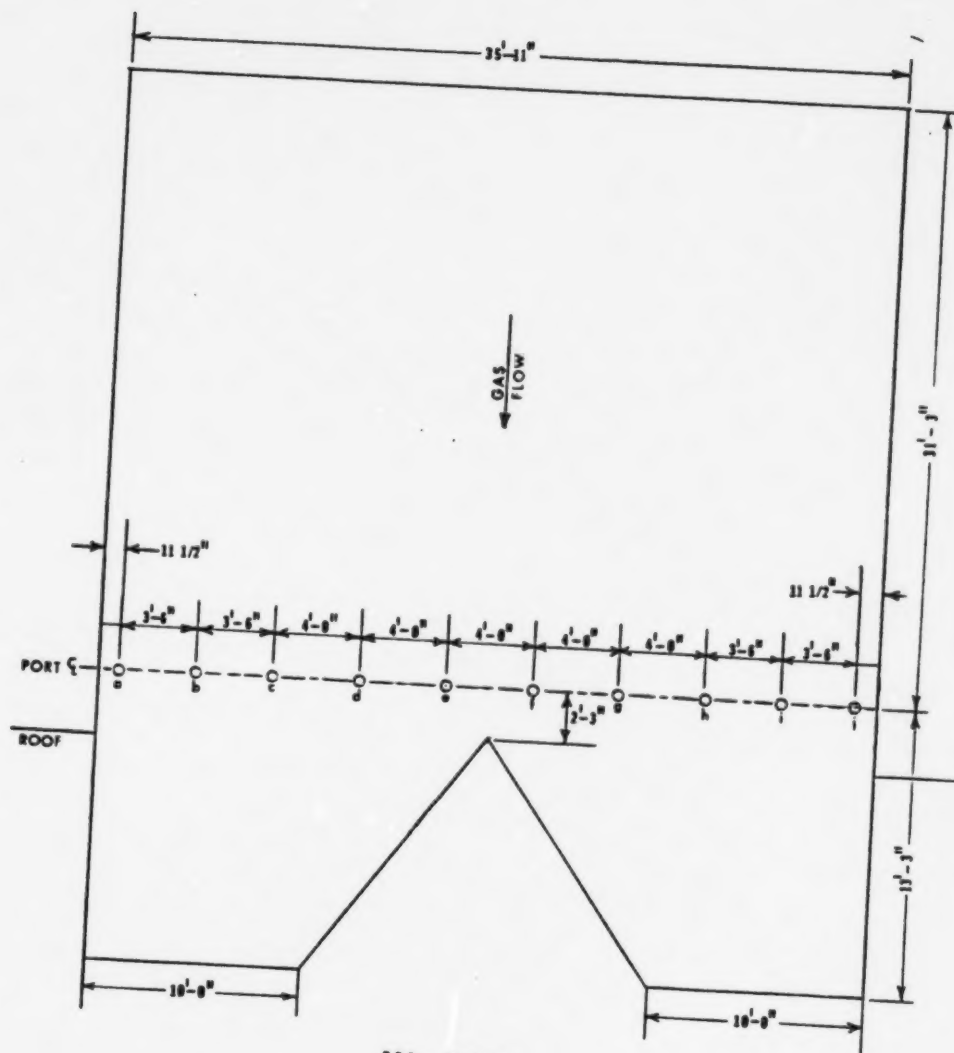


FIGURE 2
ESP OUTLET
(CROSS SECTION)



ESP OUTLET
(DUCT A & B SAME)
FIGURE 3

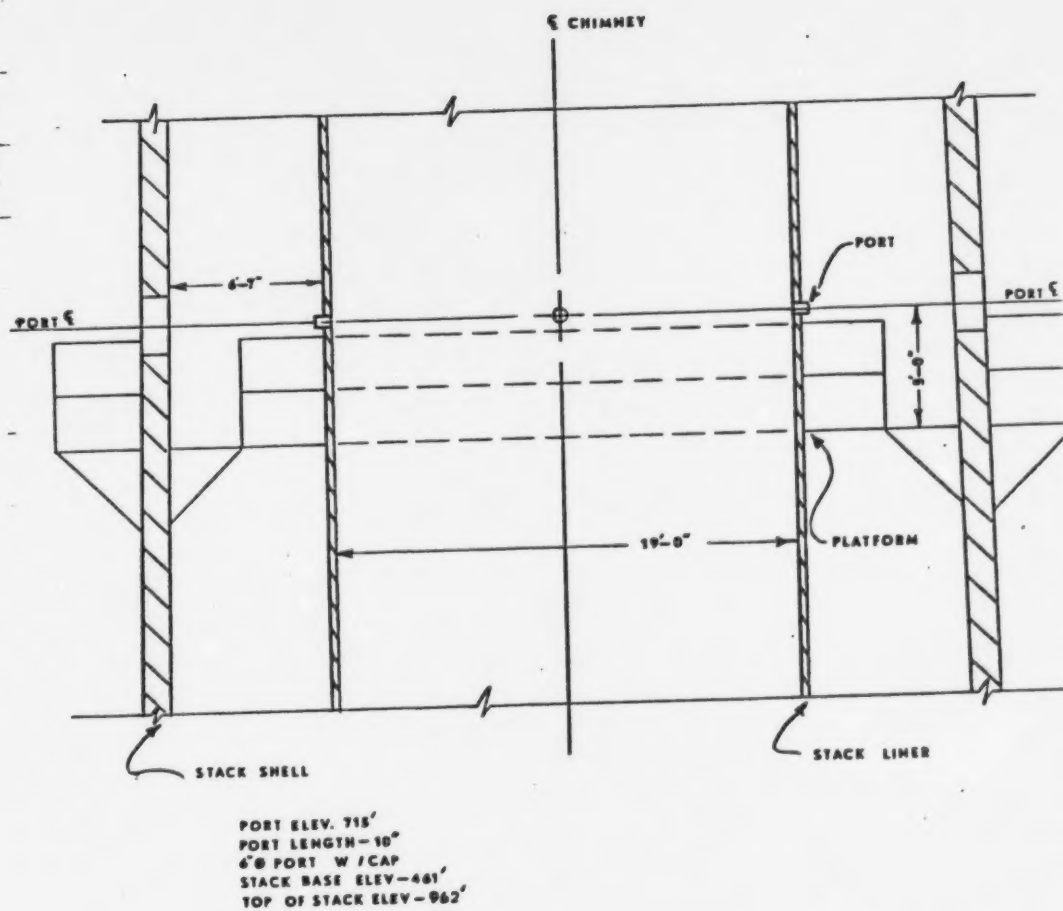
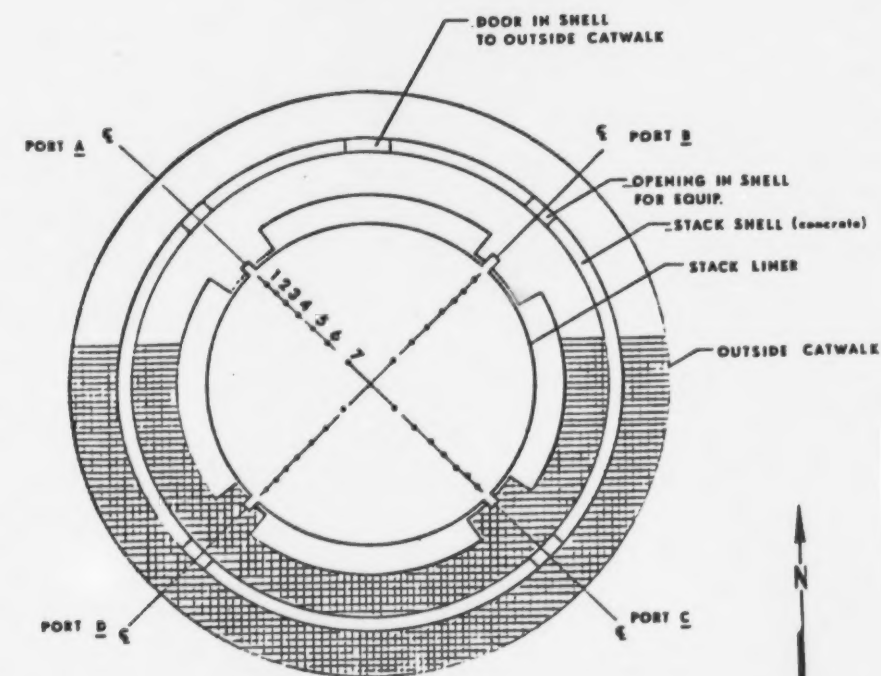


FIGURE 4
 STACK
 SIDE VIEW - SECTION



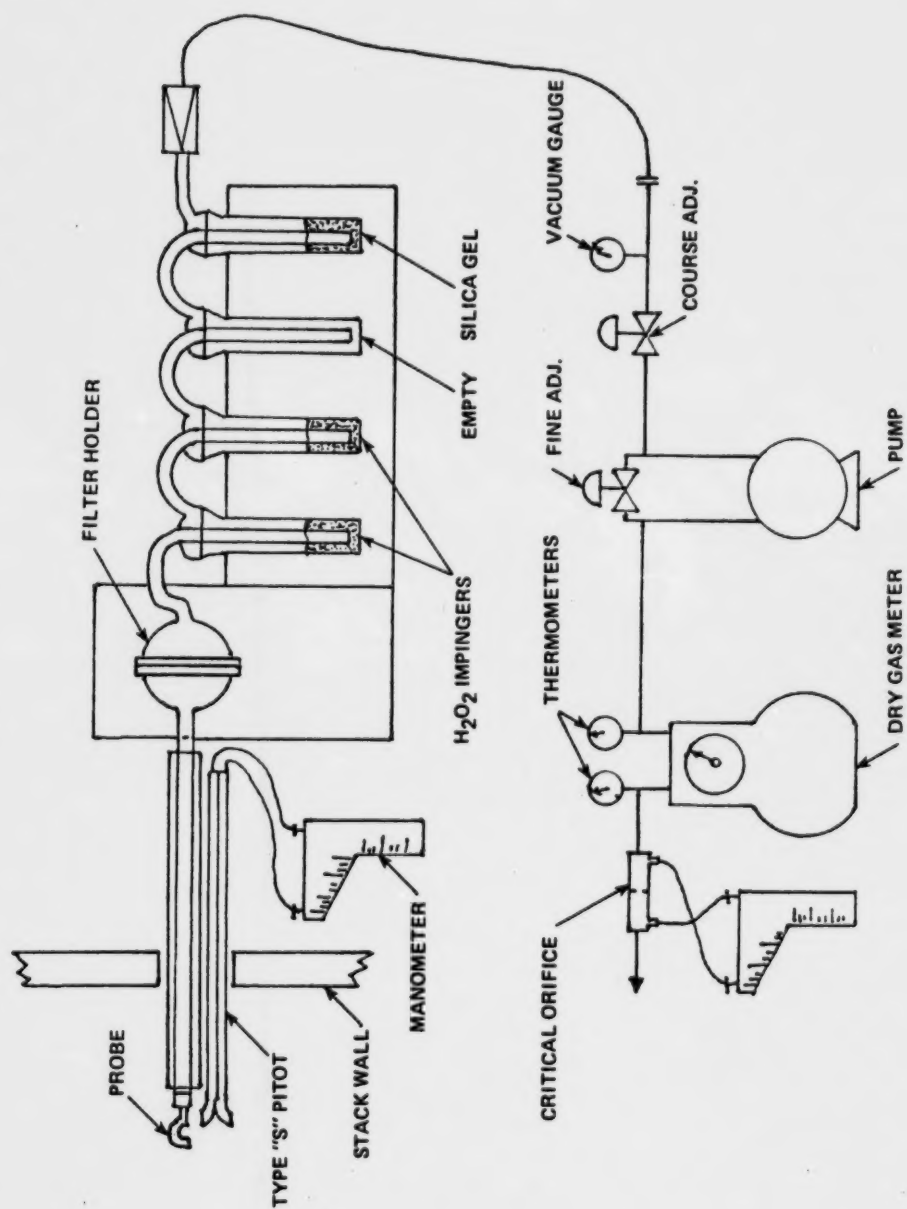
CATWALK INSIDE-OUTSIDE SHELL
19'-0" I.D. LINER AT PORT ELEV.
TOP OF BREACHING ELEV. 493'

PORT ELEV.-715'
PORT LENGTH-10"
6 @ PORT W / CAP

DISTANCE FROM OUTSIDE PORT TO POINT (IN.)			
1. 14 1/8	3. 32 5/16	5. 85 13/16	7. 93 7/16
2. 23	4. 43 5/16	6. 71 5/16	

FIGURE 5
STACK
CROSS-SECTION

FIGURE 6



the sample box contained an ice bath with four glass impingers connected in series by glass u-tubes. The first impinger contained 100 ml. of 80% isopropanol. The second and third impingers contained 100 ml. of 3% or 12% hydrogen peroxide for the stack and ESP, respectively. The final impinger contained approximately 200 gm. of indicating silica gel. The ice bath-impinger combination collected the SO₂ and cooled and dried the flue gases sampled to prevent damage to either the pump or dry gas meter. It also allowed a determination of the stack gas moisture content for each run.

Heated stainless steel (Lear-Siegler) liner probes were used at all locations. The probes were horizontal for sampling at all locations. A 10-foot probe was used at the ESP outlets and an 8-foot probe was used at the stack. Type K thermocouples attached to the probes were used to measure temperature. A close visual inspection of the sample tips and "s" pitots was made after each run. Stack-duct pressure was measured using a stainless steel tube attached to a water column u-tube manometer. Random ports were sampled.

A Taylor aneroid barometer, (I.D. No. 102 - calibrated against a mercury column barometer) located in the Kenvirons laboratory trailer, was used each day to determine barometric pressure. The difference in elevation between the lab and each test location was determined and a correction made based on 0.11 inch mercury less pressure per 100 feet elevation.

Reeve Angel (900 AF) fiberglass filters were used for particulate collection. The glass filter holders were maintained at $248^{\circ} \pm 25^{\circ}\text{F}$. The filters had been tared and stored in new plastic petri dishes with tops pursuant to method 5 (see Appendix B for filter log).

As required, the pitot systems and the sampling trains were leak-checked before and after each test run. In all instances, the vacuum on the pitot system remained stable for at least fifteen seconds at 3" H₂O (or more) and the sampling train leaked less than 0.02 CFM at a vacuum of 15" Hg in pretest check and at the maximum sample vacuum during post-test check. Leak checks were made through the probe tip.

Moisture content, temperatures, and flowrates from previous test data were used to set sample conditions for Run 1. Subsequent runs were based on acquired data.

As a part of our quality assurance program and pursuant to method specifications the following equipment items were calibrated prior to and after testing:

1. console dry gas meter
2. console meter inlet/outlet thermometers or thermocouples
3. console timer
4. in stack-duct thermocouple

5. sample box hot side bimetallic thermometer
6. sample box cold side bimetallic thermometer
7. tip inside dimensions
8. "s" pitot for dimensions

We participated in the EPA interlaboratory QA program during November, 1979, wherein all three console dry gas meters were compared with a standard calibrated orifice provided by EPA. As noted in Appendix F, the units were off by -1.73% (D-328), -0.43% (D-467) and -0.83% (D-494), which is well within the mean value of all participants (-2.1%). All calibration data are presented in Appendix F.

We also participated in another EPA interlaboratory QA program during August, 1980 wherein all three console dry gas meters were compared with a standard calibrated orifice provided by EPA. As noted in Appendix F, the units were off by 0.2% (D-328), -0.03% (D-467) and 0.4% (D-494), which is well within the mean value of all participants (-2.1%).

All equipment was prepared and set up each day and process conditions monitored until all systems were "go". After receiving the Bechtel - EPA - L G & E "go" for a test, all three teams would start sampling. Table 5 presents a summary of sample times, teams, port-point information, and sample code data. As may be noted, the sample period was more or less simultaneous. If one team encountered shutdowns the other teams also shutdown and waited to resume sampling.

After the sample was collected, the trains were purged for 15 minutes per Method 8. Final leak checks were then performed and then the probes and sample boxes were transported to the lab for sample recovery.

3.2.2 Sample Recovery and Analysis

3.2.2.1 Particulates

Kenvirons, Inc. field laboratory (36-foot long trailer) was used for all sample recovery operations. A quasi-clean area was maintained for glassware cleanup and probe rinsing. A 12-18-foot-long (6-foot sections) stainless steel rod with a nylon brush attached was used to clean the probes. Per the method, six penetration cycles were used while spraying reagent grade acetone from a glass Guth wash bottle. The rinse sample from the probe was captured in a glass funnel into a glass sample container with a teflon cap liner.

A small nylon brush was used to remove the sample and clean the sample tip and glass tube up to the filter using rinse acetone. The particulate rinse sample was recovered using the "standard" (Method 5) acetone rinse. Blanks of acetone, 80% IPA, 12% H₂O₂, and 3% H₂O₂ were taken for each run. All blanks and samples were given a special laboratory number, as noted in Table 5. Filters also had a second Kenvirons number.

GEORG[NATION CODE (CONTINUED)]

Can I use

11/10/2001

Official Start Time

1000

No. Ports

Minutes Per Point

SAMPLE DATA-LAB. CH

Acetone

Levy from the

Net Weight

 (H_2O_2) Final Mo (H_2O_2) F (mol) Me

Imp. No. 4 Prel. Me

Total Moisture (M/G)

1% H_2O_2 Blank Cont.

RELATIVE FIELD NO.

20% concentration (1)

Average ΔH (in. H₂O)

.....

1

[illegible]

The volume of condensate, etc., in each impinger after the run, was determined by weighing the impinger plus contents to the nearest 0.1 gram. Since we had the tare weight, the difference in weight was assumed to be moisture condensate from which the stack-duct moisture content was determined.

The 4-inch fiberglass filter was carefully removed from the glass holder and placed into the original petri container (plastic). Laboratory tweezers were used to scrape the silicone rubber gasket to remove all traces of the filter. High SO_2 concentrations at the stack and ESP caused deterioration in the red-color 4-inch silicone gasket which was carefully prevented from being added to the sample filter container. It was also noticed that a red-color film had collected on the first impinger insert and it was very hard to remove. This film was thought to be coming from the red silicone gasket. This red film was noticed during sample recovery through out the ten-day test period. New silicone gaskets were used for each run. The acetone rinse and blank samples plus the filters were returned to the Kenvirons laboratory in Frankfort for analysis.

Upon return to the Frankfort laboratory the filters were desiccated and weighed to constant weight as noted in Appendix B. The acetone samples and blanks were measured volumetrically then evaporated in tared beakers (glass) desiccated and weighed per method 5 procedures. The weight gain by each filter was added to the corresponding acetone rinse weight (less blank correction) and reported as total particulate weight for that test (Appendix B).

Due to a malfunctioning valve in one of the laboratory desiccators, the acetone evaporations from Day 2 may have been cross-contaminated, and some particulate lost in the desiccator. The ESP-B-2 acetone rinse beaker was closest to the valve. Beakers ESP-A-2 acetone rinse and ESP-B-2 acetone blank were the only 2 beakers immediately adjacent to ESP-B-2 beaker. They received some loose dust from the ESP-B-2 spill. This loose dust, however, was deposited on top of firmly caked particulate matter in ESP-A-2 acetone rinse beaker and could be removed simply by pouring the loose dust back into ESP-B-2 acetone rinse beaker; loose dust also fell into ESP-B-2 acetone blank (empty) and was returned to ESP-B-2 acetone rinse in the same manner. The remainder of the loose dust was deposited within the desiccator itself and was returned to ESP-B-2 acetone rinse beaker using a camel hair brush and tweezers, although some particulate loss did occur.

3.2.2.2 SO_2

The SO_2 sample recovery operation consisted of first checking all impingers and the in-line filter for signs of transfer or backflush of either the IPA or H_2O_2 . Apparently high gas temperature and/or high duct vacuum caused one EPA evaporation (impinger No. 1 IPA loss of weight) for run ESP-B-7 as noted on Table 5.

Each impinger was weighed (Ohaus 710 triple beam balance) and the weight noted to determine moisture content. Next, the IPA was discarded and the contents of the second and third impingers were poured into a 1000 ml.

graduated cylinder. Afterwards, these two impingers and all connecting glassware tubes between them were rinsed twice with deionized distilled (D-D) water, and the washings added to the 1000 ml. cylinder. The SO₂ sample recovery was completed by diluting the contents of the cylinder to exactly 1000 ml. with D-D water. The sample was then sealed in a sample container (glass, teflon liner cap).

A preliminary sample analysis (field titration) was performed on all samples for SO₂. These values were compared with Draeger tube grab samples also taken during the runs to establish approximate sampling representation accuracy (see Table 5).

Final sample analysis was performed at the Frankfort laboratory by first checking for sample loss and then pipetting a 10 ml. aliquot from the 1000 ml. sample container into a 250 ml. or 500 ml. Erlenmeyer flask, then adding 40 ml. of 100% isopropanol and two to four drops of thorin indicator. This solution was titrated to a peach-pink end-point using a previously standardized 0.0100 N (± 0.0002 N) barium chloride. The titrations were repeated and each time the replicate titration agreed within one percent or 0.2 ml.

Standardization of the 0.0100 N (± 0.0002 N) barium chloride was performed at least once per day against 25 ml. of standardized sulfuric acid to which 100 ml. of 100% isopropanol had been added. For test runs one through eight, the sulfuric acid had been previously standardized in Kenvirons Frankfort laboratory using 0.0100N (± 0.0002 N) NaOH which had been standardized against a potassium acid phthalate primary standard.

Due to the primary purpose of the scrubber demonstration program being SO₂ control technology, an emphasis was placed on accuracy in SO₂ measurements at all locations. Several cross-check measurements were thus made at several locations. L G & E operated a continuous Lear Siegler (LSI) SO₂ monitor at the stack location and Dupont instruments prior to and downstream from the scrubber. After the first few days of testing, stack SO₂ values did not agree well enough. L G & E had the LSI recalibrated against a 101 PPM SO₂ tank provided by Kenvirons. After recalibration, the Kenvirons values were still too high by 10-20% (see Table 5 preliminary day 1-8 versus Table 2).

At the end of the eighth day of the testing program it was discovered that the normality of the standard sulfuric acid used to determine the barium chloride normality had changed. Mold growth had apparently decreased the normality of the acid by fifteen percent. This would have effectively increased (thus our values were too high) the measurement of the stack and ESP sulfur dioxide concentration by fifteen percent. On day nine and ten a new commercially prepared (Fisher Scientific) solution of $.02 \pm .0002$ normal sulfuric acid (lot #704099) was used to determine the normality of the barium chloride. This solution contained a mold inhibitor. The same standard sulfuric acid solution was used to determine the barium chloride normality for the final sample titrations performed in the lab and also for the Kenvirons, September, 1980, sulfur dioxide determinations of EPA interlaboratory quality assurance standard sample. The EPA interlaboratory quality assurance results for September, 1980, Performance Audit Program Method 6 (SO₂), are noted in Appendix F. Blanks were also run with each series of

samples by titrating a 10 ml. aliquot from both the 3% and 12% H₂O₂ solutions that were freshly prepared for that sampling day. The H₂O₂ blanks were titrated using the same procedure used for sample analysis.

The results of the sample and blank analysis are presented in Appendix C.

In addition to sulfuric acid standardization, the reagent grade isopropanol was checked for peroxide impurity in Kenvirons' Frankfort laboratory using the procedures given in EPA Method 8.

3.2.3 Carrier Gas Sampling and Analysis

An integrated EPA Method 3 gas sample was taken simultaneously (same probe) with the Method 5-8 sample at the stack for each run. Figures 7 and 8 illustrate the Method 3 sample train and the probe design. The sample was collected in a 100-liter Tedlar bag which was analyzed by Orsat in the field after each run. The bag was leak checked per Method 3. Several 100 ml. Orsat analyses were performed to obtain a repeatable value.

Carrier gas sampling at all locations also was performed using a Fyrite kit (CO₂ and O₂ only). Several samples were taken during each run from random ports and the results recorded. The average of all measurements was used for run calculations. A stainless steel tube was inserted into the duct several feet beyond the port wall to obtain the sample. The kit was purged pursuant to instructions (Ambac and Bacharach).

For data reduction, Orsat results were used for stack calculations and average Fyrite results were used for the ESP A and B calculations.

3.3 Daily Log Discussion

The following presentation identifies testing procedure problems, plan changes, and corrective action that should be considered in evaluating data and results. Activities during each onsite day are discussed.

7-14-80

- * Rick Melloan and Dave Watson reported boiler problems and consequent testing delay of several days.

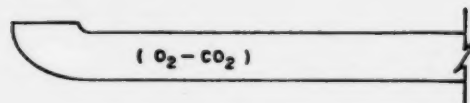
7-16-80

- * Dave Watson reported boiler problems resolved, requested testing commence 7-17-80.
- * Equipment set-up, final preparations, leak checks
- * Stack elevator problem and inability to use bullet hoist for equipment transport noted.

7-17-80 (Run 1)

- * Barrel hoist in stack made temporarily operable

GAS
FLOW



SIDE VIEW

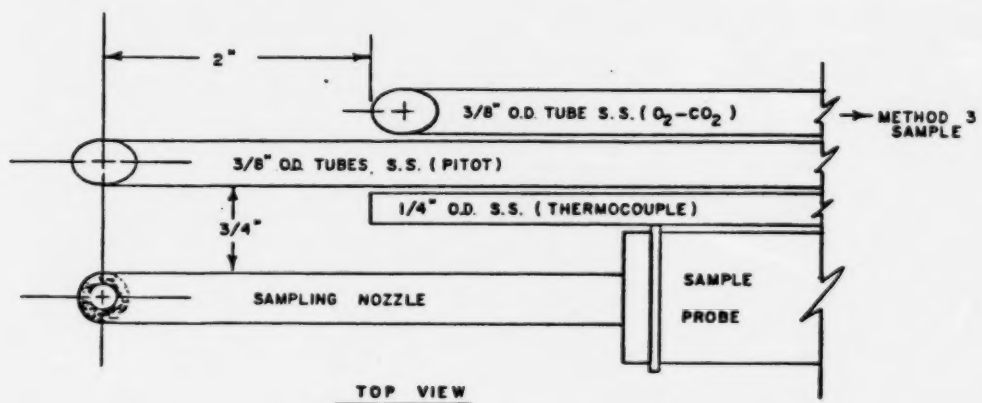


FIGURE 7

EPA METHOD 3 PROBE ATTACHED TO METHOD 5 PROBE

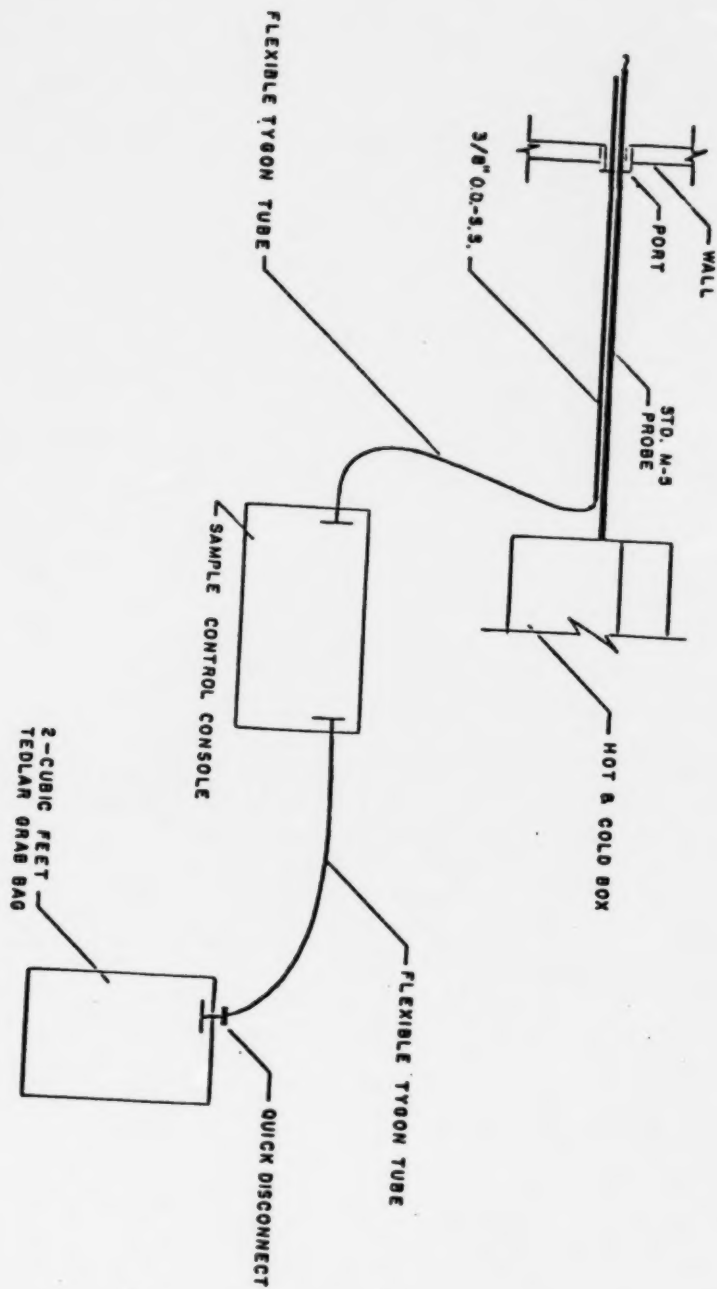


FIGURE 8
 INTEGRATED GAS SAMPLING TRAIN
 METHOD-3

- * IPA backup in impingers on one of the ESP sampling units (due to ambient heat) wet a hot filter, necessitating filter change and reweighing.
- * 8-ft. probe at stack had short in heater and a crossthreaded union - repaired by 12:30 p.m.
- * 12:30 p.m. coordination meeting. Testing to start at 2 p.m. Scrubber reported tuned and operating.
- * 2:00 p.m. - Probe tip at stack replaced with smaller tip due to higher flow rates.
- * Stack team commences sampling at 2:15 p.m.; ESP starts at 2:30 p.m.
- * 3:00 p.m. - Load at 240 MW; LSI monitor reads 70-80 ppm SO₂ at stack, 2400 ppm SO₂ at ESP outlet.
- * Testing finished at 5:45 p.m.; cleanup commenced.
- * 6:20 p.m. - Dave Watson requests no sulfate cleanup by Kenvirons
- * Completed cleanup at 8:20 p.m.; titrations begun before leaving.

7-18-80 (Run 2)

- * Told at 10:00 a.m. to begin testing when we are ready
- * Leak check problems at stack corrected by 10:30 a.m.
- * Testing began at 11:00 a.m.
- * 11:25 a.m. - Stack SO₂ at 70 ppm; ESP Outlet SO₂ at 2200 ppm.
- * 1:40 p.m. - Stack SO₂ unchanged; ESP Outlet has risen to 2300 ppm SO₂.
- * Testing completed by 2:30 p.m.; sample recovery and preliminary titrations completed by 7:00 p.m.

7-19-80 (Run 3)

- * 15% moisture at stack during Run 2 reported; were informed by Bud Carter that ESP-B reheat had tripped off 2 or 3 times.
- * Testing began at 10:33 a.m. after leak check problems at ESP-A had been corrected.
- * Stack temperature fluctuating between 140° - 190°F at noon, per Kenvirons instruments (control room reports LSI readings of 150 - 151°F).
- * ESP-B reported to have lost reheat from 11:50 - 11:54 a.m.
- * Testing completed by 2:00 p.m.
- * Red color on wet filter noted during cleanup

7-20-80 (Run 4)

- * Unit 5 scrubber washing at 10:00 a.m. causes high ambient SO₂ at ESP, delaying test startup.
- * Both the LSI monitor and Kenvirons' Draeger tubes check within 25 ppm on 2 tanks having 300 ppm and 600 ppm SO₂; yet, LSI still reads 70-90 ppm SO₂ in stack while Draeger tubes read 150 - 200 ppm. Dave Watson reports 15% stack moisture is reasonable.

- * Control room reports at 12:25 p.m. 100 ppm SO₂, 150°F, and 6.8% O₂ at stack; at ESP outlet, SO₂ between 2400 - 2600 ppm.
- * Testing commenced at 12:30 p.m.
- * Testing completed by 3:00 p.m. ESP obtained only one good Fyrite measurement at each side before last of chemicals expired.

7-21-80 (Day 5)

- * Began testing at 10:30 a.m.
- * Control room at 10:35 a.m. reported 80 ppm SO₂, 140°F, and 7.2% O₂ at stack, 2400 - 2500 ppm SO₂ at ESP outlets.
- * Fyrite chemicals for O₂ replenished via courier at 11:30 a.m.
- * Testing completed by 2:00 p.m.
- * Dave Watson reiterated problem with discrepancies in stack SO₂ values, mentioned possibility of engaging a third party to perform M6 samples.
- * Pink filters again reported, particularly at ESP; suggested explanation is breakdown of silicone gasket on hot box by heat and SO₂.

7-22-80 (Day 6)

- * Equipment setup at stack delayed until 10:30 a.m. by elevator repairs.
- * ESP-B reheat reported off at 9:30 a.m.
- * Testing OK'd at 10:30 a.m., commenced at 10:45 a.m.
- * Mostardi-Platt team arrived to double-check stack SO₂ readings by M6.
- * ESP-B reheat revised and coming back up; hence, expect less stack stratification.
- * ESP-B reading erratic ΔP at noon; reported condensate in pitot lines and tried to correct problem by blowing out lines, with only partial success.
- * Control room at 12:40 p.m. reported 130 - 150 ppm SO₂, 150°F, and 6.8% O₂ at stack; 2400 - 2500 ppm SO₂ at ESP outlets.
- * Stack team at 12:45 p.m. reported 200 ppm SO₂ per Draeger tubes.
- * Testing completed by 3:00 p.m.
- * Dave Watson reported at 3:00 p.m. that L G & E's pH probe had malfunctioned, causing loss of full control, in turn possibly causing high SO₂.
- * Dave Watson and Mostardi-Platt team reported M6 readings of stack SO₂ to be 160 - 190 ppm, whereas Kenvirons' Draeger tubes indicated 200 ppm and the LSI monitor only 90 ppm, thus supporting the Kenvirons stack data.

7-23-80 (Day 7)

- * Dave Watson reported at 8:30 a.m. a problem reaching full load (now at 150 - 160 MW), requested use of Kenvirons' 100 ppm SO₂ tank for low-scale calibration of LSI Stack SO₂ monitor (tank delivered at 1:00 p.m. this day).

- * Bud Carter at 9:15 a.m. reported wet coal was delaying attainment of full load.
- * Mickey Martin at 12:10 p.m. reported load down to 200 MW.
- * Control room reported 240 MW load at 2:15 p.m.
- * Fire in L G & E Mill at 2:55 p.m. brought load down to 220 MW by 3:00 p.m.
- * LSI technician acknowledges problems with its stack SO₂ monitor in the low range; N₂ injections show a 55 ppm reading should be 0 ppm.
- * Mickey Martin reported at 3:30 p.m. that load was still 200 - 220 MW, that today would have to be scrubbed as an official Acceptance Test day, but that Kenvirons should to ahead and test anyway.
- * Testing commenced at 4:20 p.m.; load at 225 MW.
- * Control room at 6:10 p.m. reported 225 MW load, 7% O₂ at stack, and 2200 ppm SO₂ at ESP outlets.
- * Sampling completed by 7:20 p.m.

7-24-80 (Day 8)

- * Stack reported leak check problems at 10:10 a.m. caused by cracked brass fitting connecting cold sample box and umbilical; brass fitting replaced
- * Began testing at 11:00 a.m.
- * Control room at 11:05 a.m. reported 92 ppm SO₂ (per LSI), 150°F, and 6½% O₂ at stack; 2100 ppm SO₂ at ESP outlets; load at 240 MW. Also reported that ESP-B reheat had been out all morning.
- * Control room at 12:50 p.m. reported 91 ppm SO₂ (per LSI), 150°F, and 6½% O₂ at stack; 2100 ppm SO₂ at ESP outlets. ESP-B reheat still out.
- * Testing completed by 1:45 p.m.
- * Mostardi-Platt results from 3 samples averaged 120 ppm SO₂ wet, 140 ppm dry; Kenvirons dry SO₂ result was 150 ppm, moisture was 15%.

7-25-80 (Day 9)

- * New standard solution prepared last night to replace spent original; comparison revealed 14% difference in accuracy explaining some of the earlier-mentioned high SO₂ values. Subsequent check against the new Fisher standard solutions revealed our original 0.0102 N standard solution to be in error by a factor of 0.85 (i.e., our standard for Runs 1-8 was 15% too high). On Days 9 and 10, we used the Fisher .02N standard.
- * Elevator repairs at 9:15 p.m. were causing stack transport/testing delay.
- * Dave Watson at 10:05 a.m. reported an hour's delay due to pH problems and high SO₂ (per LSI); offered us a supply of Bechtel's 0.02 N standard solution.
- * Testing began at 11:45 a.m.
- * Control room at 1:40 p.m. reported 95 ppm SO₂ (per LSI), 150°F, and 7% O₂ at stack; 2500 - 2700 ppm SO₂ at ESP outlets; load at 240 MW; and ESP-B reheat on but still low.
- * Testing completed by 2:25 p.m.

7-26-80 (Day 10)

- * Control room at 8:40 a.m. reported 158 ppm SO₂ (per LSI), 155°F, and 7½% O₂ at the stack; 2300 - 2500 ppm SO₂ at ESP outlet; load at 240 MW as of 8:38 a.m.; and ESP-B reheat in service but low. By 9:15 a.m., the SO₂ values had changed to 137-142 ppm at the stack and 2400 - 2600 ppm at ESP outlets.
- * Testing began at 9:30 a.m.
- * Dave Watson reported that cause of LSI's low SO₂ values was that the blowback system had been destroyed by lightning.
- * Control room at 11:30 a.m. reported 135 ppm SO₂ at ESP outlets and ESP-B reheat out as of 10:53 a.m.
- * Testing completed at noon; ESP Fyrite filter found to be leaking; problem corrected and Fyrites rerun.

4.0 DATA REDUCTION

A first phase of data reduction was performed in the field following each run. The phase involved determining percent isokinetic which required O_2 and CO_2 results plus moisture condensate volume in addition to sample parameter averages (stack-duct temperature, pitot ΔP , orifice ΔH and DGM in-out temperatures). Copies of calculations performed are presented in Appendix D. Equations used were taken from the Federal Register for Method 5/8, August 18, 1977.

The average orifice ΔH for each run was used to determine a DGM correction factor " γ " which is input to Equation 1, Appendix D. A multipoint calibration curve relating these two parameters was developed for each console meter prior to testing (see Appendix F). " γ " values are presented in Table 2.

After the final particulate filter weights and rinse capture mass were determined, the second and final phase of calculation was performed. Appendix B presents analytical particulate results that were input (mg) to Appendix D to finally calculate grain loading and lb/MMBTU, etc.

After the SO_2 analysis was complete, (Appendix C) sample volumes from Appendix D (VMSTD) were transferred to determine SO_2 in lb/DSCF, lb/MMBTU, lb/hr., and ppm, as indicated in Appendix E.

The final calculation checks were made by computer for both particulate and SO_2 . Results are in Appendix F.

5.0 QUALITY ASSURANCE

The quality assurance (QA) task for this project was informal, yet consistent with program goals, priorities and budget constraints. We did not establish a formal program per se, such that third-party independent audits were performed or samples were split or spiked and sent to third party laboratories for analysis.

The QA effort primarily consisted of:

- * A comprehensive equipment pre-and post-test calibration effort,
- * Adhering to Method 5-8 requirements which have built-in checks,
- * First calculations manually followed by double checking calculations by computer,
- * Some confirmation measurements using less expensive, less accurate samplers such as a Draeger for SO₂, and
- * Comparing all measurement results with expected values based on theory or previous tests. (Abbreviated on instruction from Bechtel.)

These QA elements are consistent with overall requirements for more formal efforts as stipulated in "Quality Assurance Handbook for Air Pollution Measurement Systems", Volume III, Stationary Source Specific Methods, August, 1977.

A comprehensive equipment calibration file was maintained (Appendix F) that included

- (1) pre-and post-test for the Method 5-8 dry gas meter (DGM),
- (2) sample tips (nozzles),
- (3) thermometers,
- (4) thermocouples,
- (5) barometers,
- (6) pitots, and
- (7) interlaboratory participation.

The DGM is probably the most critical element; thus, we confirmed its accuracy through three independent means. We maintain a reference DGM that has been calibrated against a wet test meter and bell prover at the University of Kentucky Department of Mechanical Engineering. A 7-point (flow rate) pre-test calibration is performed on each Method 5 console and a curve established (see Appendix F). After a run is completed and the average orifice ΔH calculated, the DGM correction (Y) is determined (see Table 2).

Our second confirmation of the DGM accuracy consisted of participating in the EPA Interlaboratory QA study of November, 1979 (Appendix F), wherein an EPA-calibrated orifice was used on our consoles. The resulting flow rates were then compared. Our three console DGM's were well within the average accuracy (2.1%). Two of the meters averaged less than 1.0%.

The third confirmation consists of a follow-up participation in another EPA Interlaboratory QA study, September, 1980 (Appendix F). This study showed that our three console DGM's were well within the average accuracy of (-2.1%). In fact, the overall average of the three meters was 0.21%.

Appendix F also presents calibrations of nozzles (tips). Several inside diameter measurements were made and averaged per Method 5 using a micrometer. Tips were inspected during and after sampling to insure no damage. If they were damaged they were recalibrated in the field before using again.

Thermocouples and bimetallic thermometers were calibrated against mercury-in-glass thermometers in an oil bath.

We also participated in an EPA interlaboratory QA study that determined the laboratory's accuracy in SO₂ titration. Appendix F presents our most recent results (we used the same standard solution - Fisher - earlier discussed). As may be noted, all participating labs averaged 1.69 - 2.72% difference while our values ranged from -2.68% to 1.0%. Our values are thus comparatively accurate, however, slightly lower than other labs.

6.0 DISCUSSION

Table 2 presented a summary of test data and results, and Table 3 presented data averages and ranges. As discussed in Section 2.0, most measured values and results were within expected ranges. Problems encountered in field measurements, sample analysis, data reduction, etc., have been generally discussed in previous report sections. The daily log also clarified problems with specific runs for evaluation purposes.

Table 6 presents a listing of sample particulate results for each run. That portion of the sample captured on the filter versus probe rinse is illustrated. A consistent percentage capture on the filter indicates consistency in sampling and indirectly particle size. Run results that are grossly out of line may indicate a poor sample or process problems.

At the stack, the filter capture varied from 54.5% to 70.6% which is a reasonably narrow range, thus indicating that all the samples may be quite good. At the ESP-A, the filter capture varied from 28.3% to 47.0%. Most values were from 30-40%; thus, the 47.0% (Run 2) indicates potential problems. As noted in section 3.2.2.1, we experienced an analysis problem that probably explains this "outside" number. The ESP-B filter capture portion varied from 25.4% to 35.5%, which indicates good consistency.

Table 7 presents particulate and SO₂ collection efficiencies for each run based on LB/MMBTU. Run 2 indicated the lowest particulate efficiency (80.05%). All other values ranged from 85% to 91%. Our analytical problem with Run 2 could have resulted in a loss of part of the sample thus giving the lower emission rate (0.362 LB/MMBTU). However, that rate is not far out of line since several other values were noted in the 0.5 LB/MMBTU range. The key point is that the ESP outlet values are reasonably high and that the scrubber is 85 - 91% efficient in removing particulate. SO₂ removal efficiency was consistent, ranging from 89.98% to 95.04% (average 93.2%).

The following additional minor points should be considered in evaluating program results.

- * The sample probes at the stack and ESP entered certain ports at less than 5° angles due to interferences (see data forms),
- * Run 6 - ESP-B - had problems with velocity measurements on 2-4 ports due to rain causing condensate in lines - corrected by blowing lines,
- * Run 3 - ESP-A - found small piece of styrofoam in impinger - probably came from storage locker - caused higher than normal vacuum during sampling.

Due to several cross-checks that were performed by Kenviron, L G & E, and Bechtel, plus the fact that our calibrations and EPA interlaboratory participation results looked good, it appears that our results should be accurate within method expectations.

In summary, aside from the minor discussed problems, all of the data and results appear to be representative of the process operation during testing.

TABLE 6
PARTICULATE SAMPLE DISTRIBUTION EVALUATION

Run	STACK					ESP-A					ESP-B				
	Filter Wt.	% of Total Wt.	Acetone Wt.	% of Total Wt.	Total Wt.	Filter Wt.	% of Total Wt.	Acetone Wt.	% of Total Wt.	Total Wt.	Filter Wt.	% of Total Wt.	Acetone Wt.	% of Total Wt.	Total Wt.
1	56.3	70.6	23.5	29.4	79.8	221.4	35.2	407.3	64.8	628.7	264.4	27.4	698.5	72.6	962.9
2	58.5	64.4	32.4	35.6	90.9	386.3	47.0	435.8	53.0	822.1	148.4	30.1	334.1	69.9	482.5
3	61.0	63.1	35.7	36.9	96.7	442.3	37.0	752.4	63.0	1194.7	397.7	26.2	1118.2	73.8	1515.9
4	58.6	64.3	32.6	35.7	91.2	358.1	33.1	722.8	66.9	1080.9	291.5	32.4	607.9	67.6	899.4
5	60.3	66.8	30.0	33.2	90.3	432.4	30.9	964.7	69.1	1397.1	313.5	29.3	755.1	70.7	1068.6
6	54.9	60.1	36.4	39.9	91.3	379.2	31.3	833.0	68.7	1212.2	499.5	27.9	1288.7	72.1	1788.2
7	44.2	54.5	36.9	45.5	81.1	248.9	34.6	469.6	65.4	718.5	291.8	31.7	629.9	68.3	921.7
8	47.2	61.9	29.1	38.1	76.3	302.8	37.4	505.7	62.6	808.5	286.7	35.5	518.9	64.5	805.6
9	67.5	68.6	30.9	31.4	98.4	323.6	35.7	583.8	64.3	907.4	315.8	25.4	926.2	74.6	1242.0
10	54.0	62.8	32.0	37.2	86.0	344.7	28.3	872.0	71.7	1216.7	218.9	30.8	491.3	69.2	710.2

TABLE 7
COLLECTION EFFICIENCIES

Run	Particulate (lb/MMBTU)			Particulate Efficiency (%)			SO ₂ (lb/MMBTU)			SO ₂ Efficiency (%)		
	A	B	Stack	A	B	Average	A	B	Stack	A	B	Average
#1	0.532	0.712	0.0895	83.18	87.43	85.305	5.99	5.84	0.348	94.19	94.04	94.115
#2	0.659	0.362	0.932	85.86	74.25	80.055	5.47	5.26	0.442	91.88	91.60	91.74
#3	0.947	1.07	0.111	88.28	89.63	88.955	5.720	5.93	0.430	92.48	92.75	92.615
#4	0.944	0.806	0.103	89.09	87.22	88.155	6.550	7.08	0.425	93.51	94.00	93.755
#5	1.110	0.920	0.102	90.08	88.91	89.495	6.040	6.510	0.364	93.97	94.41	94.19
#6	0.990	1.490	0.102	89.70	93.15	91.425	6.260	6.540	0.641	89.76	90.20	89.98
#7	0.589	0.847	0.102	82.68	87.96	85.32	5.100	5.290	0.357	93.00	93.25	93.125
#8	0.625	0.649	0.0893	85.71	86.24	85.975	5.330	5.190	0.335	93.71	93.55	93.63
#9	0.789	1.200	0.110	86.06	90.83	88.445	6.930	7.150	0.349	94.96	95.12	95.04
#10	0.962	0.593	0.102	89.40	82.80	86.10	5.900	5.890	0.359	93.92	93.90	93.91

Appendix E. Field Data Forms

(Sequence: ESP-A-(1-10); ESP-B-(1-10); and STK-(1-10))
(Start on pp E-1; E-94; and E-187)

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC.
PARTICULATE EMISSION TEST

PROJECT NUMBER 80019
FACILITY L.G.E.
RUN NUMBER BEC-ESP-A-1

FIELD DATES 7-17-80/7-26-80
SAMPLING LOCATION ESP-OUTLET-A
SAMPLING DATE 17 July 1980

FIELD CREW

PROJECT CHIEF R.C. Tussey Jr.
TESTING ENGINEER 1 V.A. LAURITZEN
2
ENGR. TECHNICIAN 1 J.J. BRALLEY
2
LAB TECHNICIAN 1
2
OTHER 1
2

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-ESP-A-1
DATE: 17 July 1980

RECORDED BY: TAL
ASSISTED BY: _____

AMBIENT TEMPERATURE: 95 °F.
BAROMETER PRESSURE: 29.66 in. Hg.
READING TIME: 0708 RECORDED BY: TAL
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-80
CORRECTED BAROMETER PRESSURE: (@ console location) 29.61
CONSOLE I.D.: D-494
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: X
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: IX IMPACT: A side STATIC: B side
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: 21 (Kenviros Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: 0.238
UMBILICAL I.D.: Silver LENGTH: 50 FT.

COMMENTS: _____

RUN NO REC-ESP-A-1
DATE 17 July 1980

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY vol
ASSISTED BY

I. PROBE ASSEMBLY

A. PROBE LENGTH 10 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER IX
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.238 IN.
B. NOZZLE IDENT. NO. 231
C. UMBILICAL LENGTH 50 FT.
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79149

III. IMPINGER CONTENT

		NORMAL		USE		FINAL AMOUNT	NET GAIN
NUMBER	TYPE*	MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	521.9 g	522.0 g	0.1
2	M	DW	100ML	12 % H ₂ O	554.3 g	602.6 g	48.3
3	GS	DRY	————	12 % H ₂ O	572.5 g	588.8 g	16.3
4	M	ADSORBENT**	~200GM	- 200 SG	677.2	693.6 g	16.4
#4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED			81.1 g
WEIGHT WITH ADSORBENT = 677.2 GM.							

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.55 ΔH_0 (POST-TEST) —

CONSOLE NO.

D-494

Y (PRE-TEST) 1.03

Y (POST-TEST) —

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☒ THERMOCOUPLE (TYPE "K") ☐ OTHER	☐ STANDARD ☐ "S" SHAPED
READING DEVICE	☒ MANOMETER (0-10 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE <u>—</u>) ☐ OTHER	☒ PYROMETER (TYPE "K") ☐ OTHER	☐ DI MANOMETER (0-10 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE <u>—</u>) U-Tube ☒ OTHER Manometer

COMMENTS:

RUN NO. BEC-ESP-A-1

KENVIRONS, INC.

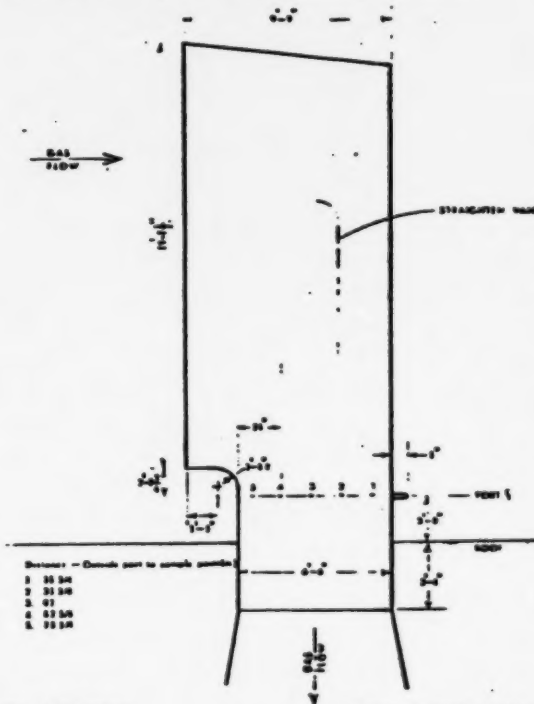
DATE 17 July 1990

STACK GEOMETRY

RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
(ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)

NOTE: SAME AS RUN _____



FOR ROUND STACKS

ESP OUTLET
(SEE SECTION 1)

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL
TO OUTSIDE OF PORT _____ IN.
B. PORT LENGTH _____ IN.
STACK DIAMETER (A-B) _____ IN.
STACK AREA: _____ IN²

A. DISTANCE FROM INSIDE FAR WALL
TO OUTSIDE OF PORT 86 IN.
B. PORT LENGTH 8 IN.
STACK DEPTH (A-B) 78 IN.
STACK WIDTH (INSIDE) 431 IN.
STACK AREA (DEPTH X WIDTH) 33618 IN²
EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN.

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
DIAMETERS

TYPE OF FLOW
OBSTRUCTION

NUMBER OF
SAMPLING POINTS

_____ DIAMETERS DOWNSTREAM OF _____

_____ DIAMETERS UPSTREAM OF _____

MIN. REQ'D. (METHOD 1)

USE

50

RUN NO. BEC-ESP-A-1
 DATE 17 July 1980

KENVIRONS, INC.
 TRAVERSE POINTS

RECORDED BY _____

POINT	PERCENT OF DIAMETER FROM INSIDE PORT TO POINT												DISTANCE FROM INSIDE PORT (IN.)	DISTANCE FROM OUTSIDE PORT (IN.)	USE (IN.)
	NUMBER OF TRAVERSE POINTS ON A DIAMETER														
	☐	☐	☒	☐	☐	☐	☐	☐	☐	☐	☐	☐			
	2	4	6	8	10	12	14	16	18	20	22	24			
1	14.6	6.7	4.4	3.2	2.6	2.1	1.8	1.6	1.4	1.3	1.1	1.1		15 3/4	
2	85.4	25.0	14.6	10.5	8.2	6.7	5.7	4.9	4.4	3.9	3.5	3.2		31 3/8	
3		75.0	29.6	19.4	14.6	11.8	9.9	8.5	7.5	6.7	6.0	5.5		47	
4		93.3	70.4	52.3	38.2	28.6	22.6	17.7	14.6	12.5	10.9	9.7	8.7	7.9	62 5/8
5			85.4	67.7	54.2	42.5	32.0	23.1	16.9	14.6	12.9	11.6	10.5		78 1/4
6			95.6	80.6	65.8	55.6	46.9	39.2	32.0	26.2	21.6	18.2	15.2		
7				89.5	77.4	64.4	53.6	44.3	36.6	29.6	24.4	20.0	16.1		
8				96.8	85.4	75.0	63.4	53.5	44.3	36.6	29.6	24.4	20.0	16.1	
9					91.8	82.3	73.1	62.5	53.5	44.3	36.6	29.6	24.4	20.0	
10					97.4	88.2	79.9	71.7	61.8	53.5	44.3	36.6	29.6	24.4	
11						93.3	85.4	78.0	70.4	61.2	53.3	44.3	36.6	29.6	
12						97.9	90.1	83.1	76.4	69.4	60.7	53.8			
13							94.3	87.5	81.2	75.0	68.5	60.2			
14							98.2	91.5	85.4	79.6	73.8	67.7			
15								95.1	89.1	83.5	78.2	72.8			
16								98.4	92.5	87.1	82.0	77.0			
17									95.6	90.3	85.4	80.6			
18									98.6	93.3	88.4	83.9			
19										96.1	91.3	86.8			
20										98.7	94.0	89.5			
21											96.5	92.1			
22											98.9	94.5			
23												96.8			
24												98.9			

COMMENTS: _____

DATE 17 July 1980

PRELIMINARY TRAVERSE

ASSISTED BY JB

[illegible]

APPROX. AVERAGE			APPROX. AVERAGE		
-----------------	--	--	-----------------	--	--

15.5 in. H₂O

100

OPENING	PRE-TRAVERSE	POST-TRAVERSE	BY
IMPACT	☐ STABLE ☐ LEAKING	☐ STABLE ☐ LEAKING	
STATIC	☐ STABLE ☐ LEAKING	☐ STABLE ☐ LEAKING	

RUN NO. REC-ESP-A-1 KENVIRONS, INC. PRELIMINARY RECORDED BY val
 DATE 17 July 1980 MOISTURE DETERMINATION ASSISTED BY _____

NOTE: SAME AS RUN _____

A. CONDENSER AND/OR SILICA GEL METHOD

BAROMETRIC PRESSURE _____ READING TIME _____
 BAROMETER LOCATION _____ READING BY _____

	CLOCK TIME	VOLUME METER(cu)	ORIFICE SETTING	METER TEMP	
				IN	OUT
FINAL					
INITIAL					
DIFFERENCE				Avg. TM	

VMSTD=17.65(Y) $\frac{VM \pm PM}{TM \pm 460}$
 = 17.65 ()
 = _____ DSGF

CONTAINER	FINAL	INITIAL	DIFFERENCE (D)	CONVERSION TO STD.
CONDENSER				D = 0.04707 = VWC
SG TUBE				D = 0.04715 = VWSG

$$\text{MOISTURE CONTENT} = \frac{(VWC + VWSG)100}{VWC + VWSG + VMSTD} = \frac{(\quad + \quad)100}{\quad + \quad} = \quad \%$$

B. PREDETERMINED VALUE

% MOISTURE 8 BASIS Previous tests

LEAK CHECK*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)		
TIME (24 HOUR CLOCK)		
BY		

* Rates < 0.02 CFM @ 15 in. Hg. vacuum are acceptable.

COMMENTS: _____

RUN NO. BEL-ESP-A-1
 DATE 17 July 1980

KENVIRONS, INC.
 NOMOGRAPH DATA

RECORDED BY val
 CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 95 °F
 BAROMETRIC PRESSURE $P_b = 29.66$
 Pbc CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 29.92) = 29.61$ IN. Hg.
 $\Delta H \theta$ REFERENCE $\Delta H = 1.55$ IN. H₂O
 TM AVG. METER TEMPERATURE = 115 °F
 % H₂O MOISTURE CONTENT = 8 % BY VOL
 PD STATIC (GUAGE) PRESSURE IN STACK = -15.5 IN. H₂O
 x 0.074
 PS ABSOLUTE PRESSURE IN STACK, $P_b - P_o = 28.96$ IN. Hg.
 PM METER PRESSURE ($\sim P_b$) = 29.61 IN. Hg.
 PS/PM = _____
 C CORRECTION FACTOR = 0.845
 TS STACK TEMPERATURE = 250 °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O
 D PROBE TIP DIAMETER = 0.238 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	250	275	300	325	350
ΔP_{REF}	0.84	0.88	0.90	~~0.94~~ 0.93	0.96

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	val	☒ STABLE ☐ LEAKING	val
STATIC	☒ STABLE ☐ LEAKING	val	☒ STABLE ☐ LEAKING	val

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	0.013 0.011 CFM @ 15" Hg.	0.005 CFM @ 7.7" Hg.
TIME (24 HOUR CLOCK)	0905 1155	1737
BY	val	val

* Rates < 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

SAMPLING

RECORDED BY val

ASSISTED BY J B

SAMPLING

SAMPLING TIME PER POINT	Z	MIN.
-------------------------	---	------

[illegible]

250/0.84 275/0.88 300/0.90 325/0.93 350/0.96

RECORDED BY 7146
ASSISTED BY J B

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 48 IN.

[illegible]

COMMENTS: _____

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80019
FACILITY L.G.E.
RUN NUMBER BEC-ESP-A-2

FIELD DATES 7-17-80 / 7-26-80
SAMPLING LOCATION ESP-OUTLET-A
SAMPLING DATE 18 July 1980

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr</u>
TESTING ENGINEER	<u>1 V.A. Lauritzen</u> <u>2</u>
ENGR. TECHNICIAN	<u>1 J.J. Bralley</u> <u>2</u>
LAB TECHNICIAN	<u>1</u> <u>2</u>
OTHER	<u>1</u> <u>2</u>

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-ESP-A-2
DATE: 19 July 1980

RECORDED BY: val
ASSISTED BY: _____

AMBIENT TEMPERATURE: 95 °F.
BAROMETER PRESSURE: 29.80 in. Hg.
READING TIME: 0735 RECORDED BY: GHR
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-80
CORRECTED BAROMETER PRESSURE: (@ console location) 29.75
CONSOLE I.D.: D-494
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: X
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: IX IMPACT: A side STATIC: R side
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: Z1 (Kenviros Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: O.238
UMBILICAL I.D.: Silver LENGTH: 50 FT.

COMMENTS: _____

RUN NO BEC-ESP-A-2
DATE 18 July 1980

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY val
ASSISTED BY T.J.B

I. PROBE ASSEMBLY

A. PROBE LENGTH 10 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER IX
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.238 IN
B. NOZZLE IDENT. NO. 231
C. UMBILICAL LENGTH 50 F
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79152

III. IMPINGER CONTENT

		NORMAL		USE		FINAL	NET
NUMBER	TYPE*	MATERIAL	AMOUNT	MATERIAL	AMOUNT	AMOUNT	GAIN
1	GS	DW	100ML	80% IPA	532.7 gm	532.9 gm	10.2 gm
2	M	DW	100ML	12 % H ₂ O	556.4 gm	608.0 gm	15.6 gm
3	GS	DRY	—	12 % H ₂ O	571.0 gm	598.1 gm	17.1 gm
4	M	ADSORBENT**	~200GM.	~200 SG	628.3 gm	647.0 gm	18.7 gm
#4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED			91.6
WEIGHT WITH ADSORBENT = 628.3 GM.							1

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: $\Delta H\theta$ (PRE-TEST) 1.55 $\Delta H\theta$ (POST-TEST) 1.03
Y (PRE-TEST) 1.03 Y (POST-TEST) 1.03

CONSOLE NO.
D-494

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☒ THERMOCOUPLE (TYPE "K") ☐ OTHER	☐ STANDARD ☐ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE) ☐ OTHER	☒ PYROMETER (TYPE "K") ☐ OTHER	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE) ☐ U-Tube ☒ OTHER <u>Manometer</u>

COMMENTS:

RUN NO. BEC-ESP-A-2

KENVIRONS, INC.

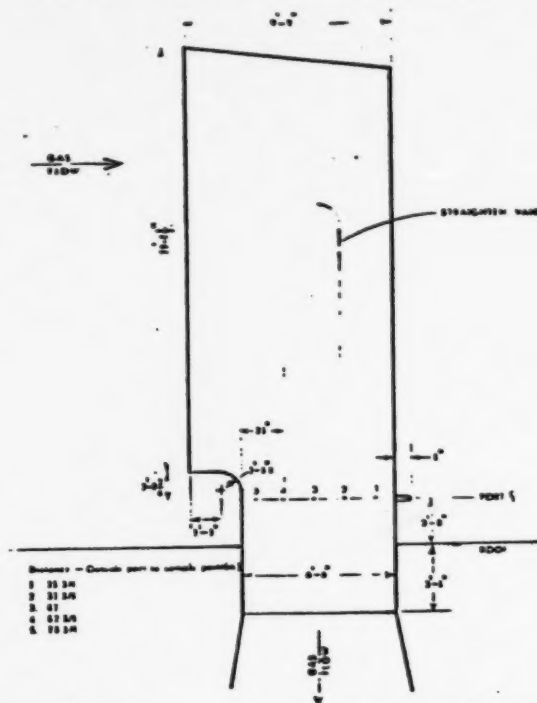
DATE 18 July 1980

STACK GEOMETRY

RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
(ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)

NOTE: SAME AS RUN _____



FOR ROUND STACKS

SEE COMMENT
SECTION 10.10.10.1

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT _____ IN	A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT <u>86</u> IN
B. PORT LENGTH _____ IN	B. PORT LENGTH <u>8</u> IN
STACK DIAMETER:(A-B) _____ IN	STACK DEPTH:(A-B) <u>78</u> IN
STACK AREA: _____ IN ²	STACK WIDTH (INSIDE) <u>431</u> IN
	STACK AREA(DEPTH X WIDTH) <u>33618</u> IN ²
	EQUIVALENT DIAMETER $(2 \frac{L \times W}{L + W})$ _____ IN

SAMPLING CROSS-SECTION IS:

NO. OF DUCT DIAMETERS _____	TYPE OF FLOW OBSTRUCTION _____	NUMBER OF SAMPLING POINTS _____
_____ DIAMETERS DOWNSTREAM OF _____		MIN. REQ'D. (METHOD 1) _____
_____ DIAMETERS UPSTREAM OF _____		USE <u>50</u>

RUN NO. BEC-ESP-A-2 KENVIRONS, INC. RECORDED BY 7726
 DATE 18 July 1980 NOMOGRAPH DATA CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 95 °F
 BAROMETRIC PRESSURE $P_b = \underline{29.80}$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 0.02) = \underline{29.75}$ IN. Hg.
 $\Delta H \theta$ REFERENCE $\Delta H = \underline{1.55}$ IN. H₂O
 TM AVG. METER TEMPERATURE = 105 °F
 % H₂O MOISTURE CONTENT = 8 % BY VOL
 P_0 STATIC (GUAGE) PRESSURE IN STACK = -15.0 IN. H₂O
 $\times 0.074 = \underline{-1.11}$ IN. Hg.
 P_S ABSOLUTE PRESSURE IN STACK, $P_b - P_0 = \underline{28.64}$ IN. Hg.
 P_M METER PRESSURE ($\sim P_b$) = 29.75 IN. Hg.
 $P_S/P_M = \underline{\hspace{2cm}}$
 C CORRECTION FACTOR = 0.83
 T_S STACK TEMPERATURE = _____ °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O
 D PROBE TIP DIAMETER = 0.238 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	250	275	300	325	350
ΔP_{REF}	0.95	0.88	0.91	0.94	0.97

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	<u>7726</u>	☒ STABLE ☐ LEAKING	<u>7726</u>
STATIC	☒ STABLE ☐ LEAKING	<u>7726</u>	☒ STABLE ☐ LEAKING	<u>7726</u>

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	<u>0.019 CFM @ 15" Hg.</u>	<u>0.011 CFM @ 8" Hg.</u>
TIME (24 HOUR CLOCK)	<u>0850</u>	<u>1350</u>
BY	<u>7726</u>	<u>7726</u>

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

KENVIRONS, INC.											
RUN NO. <u>BEC-ESP-A-2</u>			PARTICULATE			RECORDED BY <u>ral</u>					
DATE <u>14 July 1980</u>			SAMPLING			ASSISTED BY <u>J.J.B</u>					
SAMPLING TIME PER POINT <u>2</u> MIN.											
PORT POINT	CLOCK TIME	METER VOLUME (DCF)	ΔP VEL. HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
AJ-1	1100	166.28 167.75	0.42	0.85	97	96	269	270		6.8	1
AJ-2	1102	168.96	0.50	1.00	100	97	280	270		7.0	
AJ-3	1104	170.00	0.50	1.00	102	98	288	270	60	7.0	
" 4	1106	171.13	0.49	0.99	104	98	292	265	60	6.9	
" 5	1108	173.35	0.60	1.20	106	99	298	265	60	8.0	
AI-1	1130	173.37	0.39	0.80	97	96	270	255		5.9	
" 2	1132	174.96	0.47	0.96	101	98	291 292	250		6.6	
" 3	1134	175.52	0.47	0.96	104	98	304	255	58	6.6	
" 4	1136	176.68	0.47	0.96	105	99	306	250	58	6.7	
" 5	1138	177.87	0.56	1.15	107	100	306	255	59	7.6	
AH-1	1146	178.73	0.28	0.57	98	101	270	235		4.8	
" 2	1148	179.65	0.33	0.67	105	99	308	255		5.0	
" 3	1150	180.63	0.38	0.77	107	99	313	255	64	5.6	
" 4	1152	181.57	0.33	0.67	107	99	313	255	62	5.1	
" 5	1154	182.58	0.40	0.81	108	100	316	260	61	5.9	
AG-1	1200	183.23	0.15	0.31	101	99	276	250		3.3	
" 2	1202	184.07	0.29	0.59	105	100	314	250		4.8	
" 3	1204	184.96	0.30	0.60	107	100	323	250	64	4.9	
" 4	1206	185.86	0.30	0.60	108	101	322	250	62	4.9	
" 5	1208	186.85	0.38	0.77	109	101	316	255	61	5.8	
COMMENTS: 1. ANALYSIS SET FOR 300"						9.					
2.						10.					
3.						11.					
4.						12.					
5.						13.					
6.						14.					
7.						15.					
8.						16.					

350/0.65 275/0.98 304/0.91 225/0.94 350/0.97

RECORDED BY val
ASSISTED BY J. J. B

[illegible]

COMMENTS: 1. SET ANEMOGRAPH FOR 250°	9.
2. " " " 300°	10.
3. ~~" " " 350°~~	11.
4. " " " 750°	12.
5. " " " 250°	13.
6. " " " 300°	14.
7. " " " 350°	15.
8.	16.

350/1.02 275/0.98 200/0.91 125/0.94 750/0.97

RECORDED BY val

ASSISTED BY J. J. B.

SAMPLING TIME PER POINT	Z	MIN.
-------------------------	---	------

[illegible]

COMMENTS: 1. SET ALTIMETER FOR 250'	9.
2. " " " 300'	10.
3. " " " 350'	11.
4. " " " 300'	12.
5. " " " 350'	13.
6. Purged glassware @ 0150 & H form 1325 to 1415	14.
7.	15.
8.	16.

260/0.85 275/0.88 300/0.91 325/0.94 350/0.97

RECORDED BY Mal
ASSISTED BY J. J. B.

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE ~ 48 IN.

[illegible]

COMMENTS: _____

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER B0019
FACILITY L.G.E.
RUN NUMBER BEC-ESP-A-3

FIELD DATES 7-17-80 / 7-26-80
SAMPLING LOCATION ESP-OUTLET-A
SAMPLING DATE 19 July 1980

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr</u>
TESTING ENGINEER	<u>1 V.A. Lauritzen</u> <u>2</u>
ENGR TECHNICIAN	<u>1 J.J. Bralley</u> <u>2</u>
LAB TECHNICIAN	<u>1</u> <u>2</u>
OTHER	<u>1</u> <u>2</u>

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-ESP-A-3
DATE: 19 July 1980

RECORDED BY: DTL
ASSISTED BY: _____

AMBIENT TEMPERATURE: 95 °F.
BAROMETER PRESSURE: 29.86 in. Hg.
READING TIME: 0800 RECORDED BY: GR
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-80
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-494
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: I
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: IX IMPACT: A side STATIC: R side
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: Z1 (Kenvirons Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: 0.238
UMBILICAL I.D.: Silver LENGTH: 50 FT

COMMENTS: _____

RUN NO BEC-ESP-A-3
DATE 19 July 1982

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY val
ASSISTED BY TTB

I. PROBE ASSEMBLY

A. PROBE LENGTH 10 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER IX
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.238 IN.
B. NOZZLE IDENT. NO. 231
C. UMBILICAL LENGTH 50 FT.
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 28-55 79/56

III. IMPINGER CONTENT

NUMBER	TYPE*	NORMAL		USE		FINAL AMOUNT	NET GAIN
		MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	525.5 gm	538.3 gm	12.8
2	M	DW	100ML	12 % H ₂ O	552.1 gm	596.8 gm	44.7
3	GS	DRY	—	12 % H ₂ O	571.7 gm	592.0 gm	20.3
4	M	ADSORBENT**	~200GM.	~200 SG	627.7 gm	617.8 gm	20.1
#4 CONTAINER WEIGHT =				TOTAL MOISTURE ADSORBED		97.9	
WEIGHT WITH ADSORBENT =				627.7		GM.	

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.55 ΔH_0 (POST-TEST) —
Y (PRE-TEST) 1.03 Y (POST-TEST) —

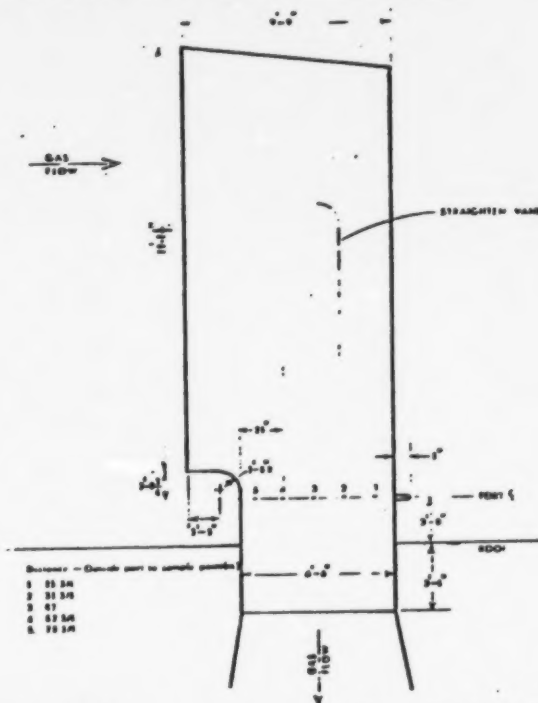
CONSOLE NO.
D-494

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☐ THERMOCOUPLE (TYPE "K") ☐ OTHER	☐ STANDARD ☐ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE) ☐ OTHER	☐ PYROMETER (TYPE "K") ☐ OTHER	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE) U-Tube ☐ OTHER <u>Manometer</u>

COMMENTS:

RUN NO. BEC-ESP-A-3 KENVIRONS, INC.
 DATE 19 July 1980 STACK GEOMETRY RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
 (ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)
 NOTE: SAME AS RUN _____



FOR ROUND STACKS

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT _____ IN.	A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT <u>86</u> IN.
B. PORT LENGTH _____ IN.	B. PORT LENGTH <u>8</u> IN.
STACK DIAMETER:(A-B) _____ IN.	STACK DEPTH:(A-B) <u>78</u> IN.
STACK AREA: _____ IN ²	STACK WIDTH (INSIDE) <u>431</u> IN.
	STACK AREA(DEPTH X WIDTH) <u>33618</u> IN. ²
	EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN.

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
DIAMETERS

TYPE OF FLOW
OBSTRUCTION

NUMBER OF
SAMPLING POINTS

_____ DIAMETERS DOWNSTREAM OF _____
 _____ DIAMETERS UPSTREAM OF _____

MIN. REQ'D.(METHOD 1) _____
 USE 50

RECORDED BY val

CHECKED BY

NOTE: SAME AS RUN

TA	AMBIENT TEMPERATURE	=	<u>95</u>	°F
Pbc	BAROMETRIC PRESSURE $P_b =$ <u>29.86</u>			
	CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 0.05) =$		<u>29.81</u>	IN. Hg.
$\Delta H \text{ ft}$	REFERENCE ΔH	=	<u>1.55</u>	IN. H ₂ O
Tm	AVG. METER TEMPERATURE	=	<u>105</u>	°F
% H ₂ O	MOISTURE CONTENT	=	<u>9</u>	% BY VOL.
PO	STATIC (GUAGE) PRESSURE IN STACK	=	<u>-1.51</u>	IN. H ₂ O
	x 0.074	=	<u>-1.12</u>	IN. Hg.
PS	ABSOLUTE PRESSURE IN STACK, $P_b - P_o$	=	<u>28.69</u>	IN. Hg.
PM	METER PRESSURE ($\sim P_b$)	=	<u>29.81</u>	IN. Hg.
PS/PM		=		
C	CORRECTION FACTOR	=	<u>0.816</u>	
TS	STACK TEMPERATURE	=		°F
ΔP	AVERAGE STACK VELOCITY HEAD	=		IN. H ₂ O
ΔP_{MAX}	MAXIMUM STACK VELOCITY HEAD	=		IN. H ₂ O
D	PROBE TIP DIAMETER	=	<u>0.238</u>	IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	250	275	300	325	350
Δ PREF	0.95		0.91		0.88

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	val.	☒ STABLE ☐ LEAKING	val.
STATIC	☒ STABLE ☐ LEAKING	val.	☒ STABLE ☐ LEAKING	val.

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	0.018 CFM @ 15" Hg.	0.016 CFM @ 15" Hg.
TIME (24 HOUR GLOCK)	1020	1312
BY	mal	mal

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

RECORDED BY nal

ASSISTED BY IJB

ASSISTED BY IJB

SAMPLING TIME PER POINT **2** MIN.

[illegible]

COMMENTS: 1. NONDISPATCH SET FOR ~~250~~³⁰⁰

9. SET NEMOGRAPH FOR 250°

2 " " " 350'

10. " " " 300'

3 " " " 300°

11.	"	"	"	350'
-----	---	---	---	------

4. ✓ " ✓ 350

12.

5. ANGLE OF PRONE MEASURED AS 4.5°

13. _____

6. MANUSCRIPT SET FOR 250°

14. _____

7. 4 11 300'

15. _____

8 " 4 " 350'

16. _____

250/245

$$300/0.91$$

350 / 293

RUN NO. <u>BEC-ESA-4-3</u>		KENVIRONS, INC.		RECORDED BY <u>val</u>							
DATE <u>19 July 1980</u>		PARTICULATE		ASSISTED BY <u>JJB</u>							
SAMPLING TIME PER POINT <u>2</u> MIN.		SAMPLING									
PORT POINT	CLOCK TIME	METER VOLUME (DCF)	ΔP VEL HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
AE-1	1134	242.30	0.07	0.15	105	104	243	280		2.2	1
" 2	1136	243.66	0.07	0.14	107	105	316	275		2.2	2
" 3	1138	243.21	0.11	0.22	~~105~~ 108	~~105~~ 108	326	270	72	2.9	
" 4	1140	243.90	0.19	0.36	111	106	337	265	72	3.6	3
" 5	1142	244.64	0.22	0.41	111	106	334	270	71	3.8	
AF-1	1148	245.20	0.10	0.22	106	105	210	275		2.8	4
" 2	1150	245.74	0.10	0.20	107	105	315	270		2.8	5
" 3	1152	246.26	0.09	0.18	107	105	323	270	80	2.6	
" 4	1154	247.02	0.24	0.49	110	105	325	265	77	4.2	
" 5	1156	247.90	0.28	0.57	112	105	327	275	70	4.8	
AG-1	1202	248.57	0.16	0.32	106	104	295	260		3.2	
" 2	1204	249.45	0.30	0.60	110	104	309	255		4.9	
" 3	1206	250.37	0.31	0.62	112	104	323	255	68	5.0	
" 4	1208	251.29	0.31	0.62	112	104	320	255	66	5.0	
" 5	1210	252.29	0.37	0.75	113	104	315	260	65	5.8	
AH-1	1216	253.28	0.33	0.72	107	103	~~251~~ 250	250		5.5	6
" 2	1218	254.27	0.37	0.75	111	103	312	250		5.9	7
" 3	1220	255.32	0.40	0.81	112	104	317	250	66	6.1	
" 4	1222	256.31	0.35	0.72	112	104	316	250	65	5.8	
" 5	1224	257.40	0.47	0.95	113	104	314	255	65	8.8	
COMMENTS: 1. <u>LET HANDS OFF FOR 250°</u>						9.					
2. " " " 300°						10.					
3. " " " 350°						11.					
4. " " " 250°						12.					
5. " " " 500°						13.					
6. " " " 250°						14.					
7. " " " 300°						15.					
8.						16.					
250/0.95						300/0.91					
350/0.98											

SAMPLING TIME PER POINT

KENVIRONS, INC.

PARTICULATE

SAMPLING

Z MIN.

RECORDED BY val

ASSISTED BY JJB

[illegible]

purged glassware @ 0.50 AN from
COMMENTS: 1. 1316 to 1333

2

3.

4.

5.

6.

7.

8.

9.

10.

11.

12.

13.

14.

15.

16.

100,

250/0.85

 $300 / 0.91$
$$350 / 0.98$$

DATE 19 July 1980

KENVIRONS, INC.
FYRITE DATA FORM

RECORDED BY val

ASSISTED BY JJA

WAS FYRITE SAMPLE TUBE PURGED BEFORE SAMPLING? ☒ YES ☐ NO

IF NO, WHAT WAS THE REASON?

WAS FYRITE ZEROED INITIALLY? ✓ YES NO

WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ✓ YES NO

WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ✓ YES NO

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE ~48 IN.

[illegible]

COMMENTS:

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80019

FACILITY L.G.E.

RUN NUMBER REC-ESP-A-4

FIELD DATES 7-17-80/7-26-80

SAMPLING LOCATION ESP-OUTLET-A

SAMPLING DATE 30 July 1980

FIELD CREW

PROJECT CHIEF R.C. Tussey Jr.

TESTING ENGINEER 1 V.A. Lawitz

2 _____

ENGR. TECHNICIAN 1 J.J. Braley

2 _____

LAB TECHNICIAN 1 _____

2 _____

OTHER 1 _____

2 _____

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-ESP-A-4
DATE: 20 July 1980

RECORDED BY: 1726
ASSISTED BY: _____

AMBIENT TEMPERATURE: 95 °F.
BAROMETER PRESSURE: 29.85 in. Hg.
READING TIME: 20-25 0745 RECORDED BY: 1726
BAROMETER I.D.: 102 CALIBRATION DATE: 6-29-80
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-494
IN TEMP. DIAL THERMOMETER I.D.: III
OUT TEMP. DIAL THERMOMETER I.D.: V
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: IX IMPACT: A side STATIC: R side
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: 21 (Kenvirons Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: 0.238
UMBILICAL I.D.: Silver LENGTH: 50 FT

COMMENTS: _____

RUN NO BEC-ESP-A-4
DATE 20 July 1940

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY ml
ASSISTED BY JTB

I. PROBE ASSEMBLY

A. PROBE LENGTH 10 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER IX
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.238 IN.
B. NOZZLE IDENT. NO. 231
C. UMBILICAL LENGTH 50 FT.
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79159

III. IMPINGER CONTENT

		NORMAL		USE		FINAL	NET
NUMBER	TYPE *	MATERIAL	AMOUNT	MATERIAL	AMOUNT	AMOUNT	GAIN
1	GS	DW	100ML	80% IPA	532.5 gm ⁵⁷⁸	542.1 gm ^{net}	20.6
2	M	DW	100ML	12 % H ₂ O	556.1 gm ⁵⁷⁸	599.5 gm ^{net}	43.4
3	GS	DRY	_____	12 % H ₂ O	571.6 gm ⁵⁷⁸	587.3 gm ^{net}	15.7
4	M	ADSORBENT**	~200GM.	~200 SG	669.1 gm ⁵⁷⁸	685.4 gm ^{net}	16.3
# 4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED		96.0	
WEIGHT WITH ADSORBENT = 669.1 GM.						215	

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.55 ΔH_0 (POST-TEST) 1.03
Y (PRE-TEST) 1.03 Y (POST-TEST) 1.03

CONSOLE NO.
D-494

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☒ THERMOCOUPLE (TYPE "K") ☐ OTHER	☐ STANDARD ☐ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-10 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE <u> </u>) ☐ OTHER	☒ PYROMETER (TYPE "K") ☐ OTHER	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE <u> </u>) U-Tube ☒ OTHER <u>Manometer</u>

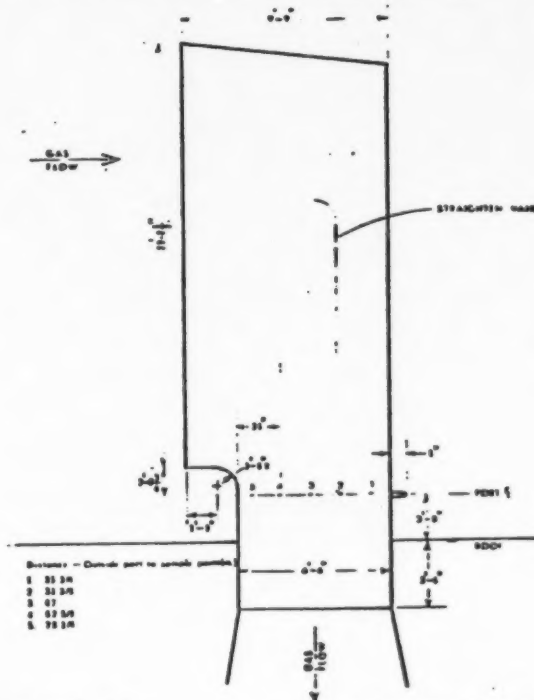
COMMENTS:

RUN NO. BEC-ESP-A-4
 DATE 20 July 1980

KENVIRONS, INC.
 STACK GEOMETRY

RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
 (ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)
 NOTE: SAME AS RUN _____



FOR ROUND STACKS

557 (001171)
 (ROUND SECTION)

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT _____ IN	A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT <u>86</u> IN
B. PORT LENGTH _____ IN	B. PORT LENGTH <u>8</u> IN
STACK DIAMETER:(A-B) _____ IN	STACK DEPTH:(A-B) <u>78</u> IN
STACK AREA: _____ IN ²	STACK WIDTH (INSIDE) <u>431</u> IN
	STACK AREA(DEPTH X WIDTH) <u>33618</u> IN ²
	EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
 DIAMETERS

TYPE OF FLOW
 OBSTRUCTION

NUMBER OF
 SAMPLING POINTS

_____ DIAMETERS DOWNSTREAM OF _____
 _____ DIAMETERS UPSTREAM OF _____

MIN. REQ'D. (METHOD 1)

USE

50

RUN NO. BEC-ESP-A-4

KENVIRONS, INC.

RECORDED BY val

DATE 20 July 1980

NOMOGRAPH DATA

CHECKED BY

NOTE: SAME AS RUN

TA AMBIENT TEMPERATURE = 95 °F
 BAROMETRIC PRESSURE $P_b = \underline{29.85}$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) = (P_b - 0.05) = \underline{29.80}$ IN. Hg.
 ΔH REFERENCE $\Delta H = \underline{1.55}$ IN. H₂O
 TM AVG. METER TEMPERATURE = 115 °F
 $\% \text{ H}_2\text{O}$ MOISTURE CONTENT = 9 % BY VOL.
 P_o STATIC (GAUGE) PRESSURE IN STACK = -14.2 IN. H₂O
 $\times 0.074 = \underline{-1.05}$ IN. Hg.
 P_s ABSOLUTE PRESSURE IN STACK, $P_b - P_o = \underline{28.75}$ IN. Hg.
 P_m METER PRESSURE ($\sim P_b$) = 29.80 IN. Hg.
 $P_s/P_m =$
 C CORRECTION FACTOR = 0.818 0.83
 T_s STACK TEMPERATURE = °F
 ΔP AVERAGE STACK VELOCITY HEAD = IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = IN. H₂O
 D PROBE TIP DIAMETER = 0.238 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	250	275	300	325	350
ΔP_{REF}	<u>0.85</u> <u>0.87</u>		<u>0.82</u> <u>0.83</u>		<u>0.87</u> <u>0.87</u>

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	<u>val</u>	☒ STABLE ☐ LEAKING	<u>val</u>
STATIC	☒ STABLE ☐ LEAKING	<u>val</u>	☒ STABLE ☐ LEAKING	<u>val</u>

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	<u>0.014 CFM @ 15" Hg</u>	<u>0.006 CFM @ 6" Hg</u>
TIME (24 HOUR CLOCK)	<u>0554</u>	<u>1500</u>
BY	<u>val</u>	<u>val</u>

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

SAMPLING TIME PER POINT 2 MIN.

44

SAMPLING TIME PER POINT	Z	MIN.
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[illegible]

COMMENTS-1. <i>NUMERICAL SET FOR 250°</i>	9.
2. " " " <i>300°</i>	10.
3. " " " <i>350°</i>	11.
4. " " " <i>250°</i>	12.
5. " " " <i>300°</i>	13.
6. " " " <i>350°</i>	14.
7.	15.
8.	16.

250/0.87 0.85 300/0.93 0.92 350/0.99 0.97

RECORDED BY val

ASSISTED BY JJB

ASSISTED BY JJB

[illegible]

COMMENTS: 1. NO MUG-RAV GOT FOR 250'

9.

2 " " " 300'

10.

3 " " " 750°

11.

4. " " " 300'

12.

5. " " " 350°

13.

6. Purged glassware @ 0.50 AM 4/10/00
1505 to 1500

14.

7.

15.

B.

16.

250/252 0.85 300/295 0.92 950/944 0.97

RECORDED BY val
ASSISTED BY TJB

WAS FYRITE ZEROED INITIALLY? ☒ YES ☐ NO
 WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ☒ YES ☐ NO
 WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ☒ YES ☐ NO

[illegible]

COMMENTS: _____

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80019

*FIELD DATES 7-17-80/7-26-80

FACILITY L.G.E.

SAMPLING LOCATION ESP-OUTLET-A

RUN NUMBER BEC-ESP-A-5

SAMPLING DATE 21 July 1980

FIELD CREW

PROJECT CHIEF R.C. Tussey Jr

TESTING ENGINEER
1 V. A. Lauritzen
2 _____

ENGR. TECHNICIAN
1 J. J. Bralley
2 _____

LAB TECHNICIAN
1 _____
2 _____

OTHER
1 _____
2 _____

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-ESP-A-5
DATE: 21 July 1980

RECORDED BY: YGL
ASSISTED BY: JTB

AMBIENT TEMPERATURE: 90 °F
BAROMETER PRESSURE: 29.85 in. Hg.
READING TIME: 0905 RECORDED BY: YGL
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-80
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-494
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: X
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: — LENGTH: —
PITOT I.D.: IX IMPACT: A side STATIC: R side
PITOT EXTENSION I.D.: — LENGTH: —
SAMPLE BOX I.D.: S-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: 21 (Kenvirons Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: O.238
UMBILICAL I.D.: Silver LENGTH: 50 FT.

COMMENTS: _____

RUN NO BEC-ESP-A-5
DATE 21 July 1980

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY val
ASSISTED BY JTB

I. PROBE ASSEMBLY

A. PROBE LENGTH 10 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER IX
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.238 IN.
B. NOZZLE IDENT. NO. 231
C. UMBILICAL LENGTH 50 FT.
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79168

III. IMPINGER CONTENT

NUMBER	TYPE*	NORMAL		USE		FINAL AMOUNT	NET GAIN
		MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	522.8 ^g	542.9 ^g	20.1
2	M	DW	100ML	12% H ₂ O	554.6 ^g	600.8 ^g	46.2
3	GS	DRY	—	12% H ₂ O	571.6 ^g	590.1 ^g	18.5
4	M	ADSORBENT**	~200GM.	~200 SG	658.4 ^g	676.5 ^g	17.7
#4 CONTAINER WEIGHT = — GM.						TOTAL MOISTURE ADSORBED	
WEIGHT WITH ADSORBENT = 658.8 GM.						102.5	

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_p (PRE-TEST) 1.55 ΔH_p (POST-TEST) —
Y (PRE-TEST) 1.03 Y (POST-TEST) —

CONSOLE NO
D-494

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☐ THERMOCOUPLE (TYPE "K") ☐ OTHER	☐ STANDARD ☐ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE —) ☐ OTHER	☐ PYROMETER (TYPE "K") ☐ OTHER	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE —) U-Tube ☒ OTHER <u>Manometer</u>

COMMENTS:

RUN NO. BEC-ESP-A-5

KENVIRONS, INC.

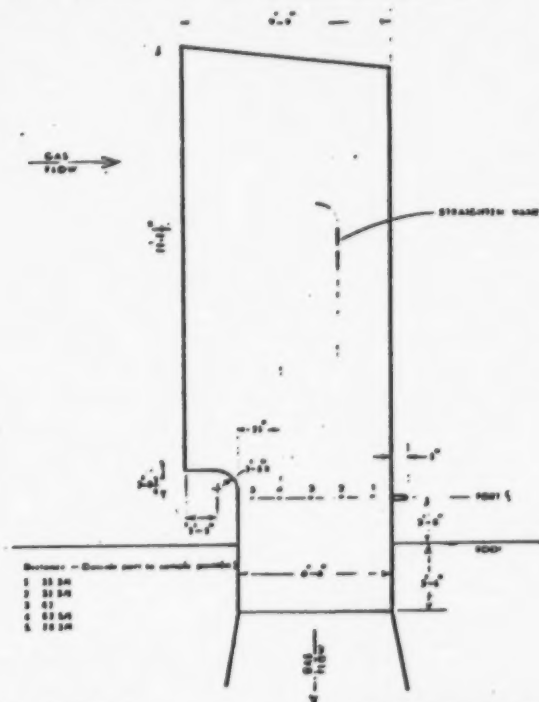
DATE 21 July 1980

STACK GEOMETRY

RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
(ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)

NOTE: SAME AS RUN _____



FOR ROUND STACKS

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL
TO OUTSIDE OF PORT _____ IN.
B. PORT LENGTH _____ IN.
STACK DIAMETER: (A-B) _____ IN.
STACK AREA: _____ IN²

A. DISTANCE FROM INSIDE FAR WALL
TO OUTSIDE OF PORT 86 IN.
B. PORT LENGTH 8 IN.
STACK DEPTH: (A-B) 78 IN.
STACK WIDTH (INSIDE) 431 IN.
STACK AREA (DEPTH X WIDTH) 33618 IN²
EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN.

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
DIAMETERS

TYPE OF FLOW
OBSTRUCTION

NUMBER OF
SAMPLING POINTS

_____ DIAMETERS DOWNSTREAM OF
_____ DIAMETERS UPSTREAM OF

MIN. REQ'D. (METHOD 1)

USE

50

KENVIRONS, INC.											
RUN NO. <u>BEC-ESP-A-5</u>				PARTICULATE				RECORDED BY <u>val</u>			
DATE <u>21 July 1980</u>				SAMPLING				ASSISTED BY <u>JTB</u>			
SAMPLING TIME PER POINT <u>2</u> MIN.											
PORT POINT	CLOCK TIME	METER VOLUME (DCF)	ΔP VEL HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
AA-1	1030	327.84	0.39	0.58	101	99	288	260		5.0	1
" 2	1032	329.80	0.41	0.82	103	100	310	265		5.7	
" 3	1034	330.80	0.37	0.68	106	101	333	275	62	5.0	2
" 4	1036	331.80	0.40	0.74	107	101	338	270	61	5.1	
" 5	1038	332.87	0.46	0.84	109	102	339	275	61	5.8	
AB-1	1044	333.80	0.30	0.64	106	102	271	270		4.7	3
" 2	1046	334.86	0.46	0.91	110	103	305	270		6.0	4
" 3	1048	335.97	0.46	0.91	111	103	330	270	60	6.0	
" 4	1050	337.01	0.41	0.76	112	104	339	270	60	5.2	5
" 5	1052	338.09	0.47	0.88	113	104	336	270	58	6.0	
AC-1	1056	338.91	0.25	0.49	108	104	275	270		4.0	6
" 2	1058	339.84	0.34	0.67	112	105	295	265		5.0	
" 3	1100	340.80	0.34	0.67	113	105	323	270	66	5.0	
" 4	1102	341.70	0.31	0.57	114	106	335	270	67	4.6	7
" 5	1104	342.63	0.35	0.64	115	106	342	270	67	5.0	
AD-1	1110	343.37	0.20	0.40	109	106	295	270		3.8	8
" 2	1112	344.20	0.25	0.49	113	106	304	270		4.1	
" 3	1114	345.06	0.27	0.53	115	107	330	270	66	4.2	
" 4	1116	345.93	0.27	0.53	116	108	331	270	66	4.2	9
" 5	1118	346.82	0.31	0.57	117	108	325	270	64	4.8	
COMMENTS: 1. NOMOGRAPH SET FOR 300"											
2. " " " 350"											
3. " " " 250"											
4. " " " 300"											
5. " " " 350"											
6. " " " 300"											
7. " " " 350"											
8. " " " 300"											
9. NOMOGRAPH SET FOR 350"											
10.											
11.											
12.											
13.											
14.											
15.											
16.											

250/0.68

300/0.94

350/1.00

KENVIRONS, INC.											
RUN NO. <u>BC-ESA-4-5</u>				PARTICULATE				RECORDED BY <u>WLB</u>			
DATE <u>21 July 1980</u>				SAMPLING				ASSISTED BY <u>JTB</u>			
SAMPLING TIME PER POINT <u>2</u> MIN.											
PORT POINT	CLOCK TIME	METER VOLUME (DCF)	ΔP VEL. HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
AE-1	1124	347.78	0.05	0.10	109	108	270	275		2.0	1
" 2	1126	347.59	0.06	0.13	110	108	276	270		2.1	
" 3	1128	348.12	0.10	0.20	112	109	313	270	72	2.8	2
" 4	1130	348.86	0.20	0.40	115	109	324	270	74	3.8	
" 5	1132	349.41	0.22	0.40	116	109	327	275	68	3.8	3
AE-1	1136	350.24	0.14	0.28	112	109	293	270		3.1	4
" 2	1138	350.87	0.14	0.28	114	110	306	270		3.1	
" 3	1140	351.38	0.09	0.18	114	110	320	270	74	2.7	
" 4	1142	352.16	0.24	0.47	117	110	319	270	68	4.1	
" 5	1144	353.01	0.27	0.53	118	110	321	270	66	4.5	
AE-1	1149	353.75	0.20	0.40	113	110	284	270		3.9	
" 2	1151	354.63	0.29	0.57	117	110	310	270		4.8	
" 3	1153	355.57	0.32	0.68	118	110	313	265	60	5.1	
" 4	1155	356.50	0.32	0.63	118	111	310	265	61	5.1	
" 5	1157	357.43	0.32	0.63	118	110	307	270	60	5.1	
AE-1	1202	358.34	0.30	0.57	114	109	291	250		5.0	
" 2	1204	359.28	0.33	0.65	117	109	292	245		5.2	
" 3	1206	360.29	0.38	0.75	118	110	306	260	60	6.0	
" 4	1208	361.25	0.33	0.65	118	110	305	275	58	5.3	
" 5	1210	362.29	0.43	0.85	119	110	303	275	60	6.8	
COMMENTS: 1. MONOGRAPH SET FOR 250°							9.				
2. " " " 300°							10.				
3. " " " 350°							11.				
4. " " " 300°							12.				
5.							13.				
6.							14.				
7.							15.				
8.							16.				
250/0.88							340/0.94		350/1.00		

RECORDED BY ral
ASSISTED BY TJB

WAS FYRITE ZEROED INITIALLY? ✓ YES NO
 WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ✓ YES NO
 WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ✓ YES NO

[illegible]**COMMENTS:**

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER B0019
FACILITY L.G.E.
RUN NUMBER BEC-ESP-A-6

FIELD DATES 7-17-80/7-26-80
SAMPLING LOCATION ESP-OUTLET-A
SAMPLING DATE 22 July 1980

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr</u>
TESTING ENGINEER	<u>1 V.A. Lauritzen</u> <u>2</u>
ENGR TECHNICIAN	<u>1 J.J. Bruney</u> <u>2</u>
LAB TECHNICIAN	<u>1</u> <u>2</u>
OTHER	<u>1</u> <u>2</u>

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: DEC-ESP-A-6
DATE: 22 July 1980

RECORDED BY: VAL
ASSISTED BY: _____

AMBIENT TEMPERATURE: 80 °F.
BAROMETER PRESSURE: 29.75 in. Hg.
READING TIME: 2300 RECORDED BY: VAL
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-70
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: 0-494
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: X
PROBE I.D.: 205 LENGTH: 10 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: TX IMPACT: A side STATIC: R side
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: 5-311
COLD BOX DIAL THERMOMETER I.D.: A - 1" DIAL
HOT BOX DIAL THERMOMETER I.D.: A - 2" DIAL
THERMOCOUPLE I.D.: Z1 (Kenvirons Shield) LENGTH: 12 FT.
NOZZLE I.D.: 231 SIZE: 0.238
UMBILICAL I.D.: Silver LENGTH: 50 FT.

COMMENTS: _____

RUN NO. BEC-STR-6 KENVIRONS, INC. RECORDED BY GHR
 DATE 7/22/86 NOMOGRAPH DATA CHECKED BY [Signature]

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 85 °F
 BAROMETRIC PRESSURE $P_b = 29.74$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 30) = 29.57$ IN. Hg.
 ΔH_R REFERENCE $\Delta H = 1.60$ IN. H₂O
 T_M AVG. METER TEMPERATURE = 105 °F
 $\% H_2O$ MOISTURE CONTENT = 15 % BY V
 P_0 STATIC (GAUGE) PRESSURE IN STACK = -0.75 IN. H₂O
 $\times 0.074 = -0.06$ IN. Hg.
 P_S ABSOLUTE PRESSURE IN STACK, $P_b - P_0 = 29.51$ IN. Hg.
 P_M METER PRESSURE ($\sim P_b$) = 29.57 IN. Hg.
 $P_S/P_M =$ _____
 C CORRECTION FACTOR = 0.785
 T_S STACK TEMPERATURE = 175 °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O
 D PROBE TIP DIAMETER = 0.2417 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	100	125	150	175	
ΔP_{REF}				3.0	

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	6:45 pm	☒ STABLE ☐ LEAKING	6:45 pm
STATIC	☒ STABLE ☐ LEAKING	6:45 pm	☒ STABLE ☐ LEAKING	6:45 pm

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	0.012 cfm @ 15" Hg	0.007 cfm @ 6" Hg
TIME (24 HOUR CLOCK)	10:12 thru 11 pm	12:23 thru 1 pm
BY	GHR + PM	GHR + PM

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

KENVIRONS, INC.
 RUN NO. BEC-STK-6 PARTICULATE
 DATE 7/22/84 SAMPLING
 SAMPLING TIME PER POINT 3 MIN.
 RECORDED BY E.H.C.
 ASSISTED BY P.H.

PORT POINT	CLOCK TIME	METER VOLUME (DCF)	Δ P VEL. HEAD (IN. H ₂ O)	Δ H ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
A-1	10:25	531.44	0.56	0.30	87	84	177	230	68	3.8	
" 2		532.91	0.65	0.39	100	85	183	225	64	4.0	1
" 3		534.75	0.85	0.52	105	85	183	225	64	5.0	1
" 4		536.09	0.85	0.52	107	85	186	225	65	5.0	1
" 5		537.23	0.80	0.48	110	86	184	225	69	5.0	
" 6		538.46	0.80	0.48	112	87	181	235	64	5.0	
A " 7	10:56	639.76	0.80	0.48	114	88	178	240	64	5.0	
D-1	10:59	640.72	0.55	0.33	107	90	179	260	68	3.8	
" 2		641.82	0.65	0.39	113	90	179	250	64	4.0	2
" 3		643.04	0.80	0.48	116	91	174	235	62	4.8	2
" 4		644.31	0.85	0.52	118	92	177	220	60	5.0	
" 5		645.52	0.80	0.48	118	92	174	200	61	4.8	
" 6		646.77	0.80	0.48	120	93	172	220	61	4.8	
D " 7	11:20	648.01	0.80	0.48	120	93	172	220	61	4.8	
C-1	11:28	648.93	0.45	0.27	111	95	169	230	68	3.4	
" 2		650.00	0.60	0.36	117	95	171	235	62	4.0	
" 3		651.13	0.70	0.42	118	95	173	240	62	4.3	3
" 4		652.33	0.75	0.45	120	95	172	240	62	4.8	3
" 5		653.50	0.80	0.48	121	95	171	240	62	5.0	
" 6		654.80	0.80	0.48	121	95	174	250	62	5.0	
C " 7	11:49	656.05	0.75	0.45	121	95	174	250	62	4.9	

COMMENTS: 1. Angle is vertical @ 45° 9.

2. Angle is vertical @ 5.0° 10.

3. Angle is vertical @ 5.0° 11.

4. 12.

5. 13.

6. 14.

7. 15.

8. 16.

KENVIRONS, INC.
 RUN NO. BEC-STK-6 CARRIER GAS RECORDED BY GR
 DATE 7-22-80 SAMPLING (ORSAT) ASSISTED BY

NOTE: SAME AS RUN

A. GRAB SAMPLE (EVACUATED BULB)

SAMPLING TRAIN DESCRIPTION:

BULB MATERIAL AND VOLUME ML

PROBE MATERIAL AND LENGTH IN

PREFILTER MATERIAL

SAMPLING LOCATION: PORT POINT

BULB NUMBER	INITIAL BULB VACUUM (IN. Hg)	VACUUM MEASUREMENT TIME (MIN.)	CLOCK TIME (24 HR. CLOCK)

B. INTEGRATED SAMPLE (TEDLAR BAG-TRAIN)

SAMPLING TRAIN DESCRIPTION:

PROBE MATERIAL AND LENGTH stainless steel 8 ft. IN.

PREFILTER MATERIAL none

PUMP TYPE diaphragm COOLING COIL stainless steel ice b

SAMPLING LOCATION: PORT A11 POINT 2-7

SAMPLING TIME: START same as before STOP

INTEGRATED SAMPLE TRAIN LEAK CHECKS*					
PRE-TEST	TIME	BY	POST-TEST	TIME	BY
☒ STABLE ☐ LEAKING	10:00	PM	☒ STABLE ☐ LEAKING	12:20	PM

* Vacuum must remain stable for 30 seconds @ 10 in. Hg.

BAG LEAK CHECK PERFORMED BY RST TIME overnight

COMMENTS: 7/21/80 - 7/22/80

KENVIRONS, INC.
DRÄGER DATA FORM

RUN NO. BEC-STK-6
DATE 7/22/80

TYPE OF DRÄGER _____

RECORDED BY 682
ASSISTED BY _____

DISTANCE FROM DUCT INSIDE WALL TO DRÄGER SAMPLE TUBE TIP - 12 IN.
DISTANCE FROM DUCT INSIDE WALL TO METHOD 6 OR 8 PROBE TIP - — IN.
SAMPLE PORT DESIGNATION C

ARE BOTH SAMPLES TAKEN FROM SAME PORT? — YES — NO

IF NO, APPROXIMATE DISTANCE FROM PROBE TIP TO SAMPLE TIP — IN.
WAS DRÄGER SAMPLE TUBE PURGED? ✓ YES — NO

IF NO, GIVE REASON FOR NOT PURGING TUBE _____

RUN NO.	TIME	N NO. OF PUMPS FOR SAMPLE	R DRÄGER TUBE READING	R X 10 N IN PPM
STK6-A	12:45	10	225	225
STK6-B	12:48	10	205	205
STK6-C	12:51	10	205	205
STK6-D	12:55	10	190	190

COMMENTS: _____

RECORDED BY PJM
ASSISTED BY _____

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 12 IN.

[illegible]

COMMENTS: _____

RUN NO. BEC-STA-6 KENVIRONS, INC.
 DATE 7-22-80 CARRIER GAS
ANALYSIS (ORSAT) ANALYZED BY PR

ORSAT LEAK CHECKS											
PRE-ANALYSIS						POST-ANALYSIS					
NO.	TIME	BURETTE LEVEL		OK	BY	NO.	TIME	BURETTE LEVEL		OK	BY
	START STOP	START STOP	START STOP				START STOP				
1	1530 1535	32.7 32.9		X	PR	1	1605 1610	42.0 42.0		✓	PR
2						2					
3						3					
4						4					

SAMPLE IDENTIFICATION	CO ₂		O ₂		CO
	READING 1	READING 2	(2-1)	READING 3	(3-2)
#6 Stack Run 6	11.0	18.8	7.8	18.8	0
	11.1	18.8	7.7	18.8	0
	11.1	18.8	7.7	18.8	0
	AVG = 11.07		AVG = 7.73		AVG = 0
	AVG =		AVG =		AVG =
	AVG =		AVG =		AVG =
	AVG =		AVG =		AVG =

CK
RS

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80019 FIELD DATES 7-17-80 / 7-26-80
FACILITY LGIE Can Run 6 Stack SAMPLING LOCATION CR6 Stack
RUN NUMBER 8 BEC-STK-7 SAMPLING DATE 7/23/80

FIELD CREW

PROJECT CHIEF R.C. Tussey Jr.
TESTING ENGINEER 1 Gary H. Revett
2
ENGR. TECHNICIAN 1 Parker H. Moore
2
LAB TECHNICIAN 1
2
OTHER 1
2

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BK-STK-7
DATE: 7-28-80

RECORDED BY: RLT
ASSISTED BY: _____

AMBIENT TEMPERATURE: 97 °F
BAROMETER PRESSURE: 29.80 in. Hg.
READING TIME: 7:41 AM RECORDED BY: RLT
BAROMETER I.D.: 102 CALIBRATION DATE: 5-29-80
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-328
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: I
PROBE I.D.: 307 LENGTH: 8 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: VII IMPACT: side B STATIC: side A
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-3112
COLD BOX DIAL THERMOMETER I.D.: C - 1" Dial
HOT BOX DIAL THERMOMETER I.D.: C - 2" Dial
THERMOCOUPLE I.D.: ZB (Kenvirons Shield) LENGTH: 9 FT.
NOZZLE I.D.: 435 SIZE: 0.241
UMBILICAL I.D.: Green LENGTH: 75 FT.

COMMENTS: _____

RUN NO REC-STX-7
DATE 7-23-80

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY 612
ASSISTED BY PM

I. PROBE ASSEMBLY

A. PROBE LENGTH 8 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER VII
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.172 IN
B. NOZZLE IDENT. NO. 435
C. UMBILICAL LENGTH 75 F
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79173

III. IMPINGER CONTENT

NUMBER	TYPE*	NORMAL		USE		FINAL AMOUNT	NET GAIN
		MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	519.5g	549.1	29.6
2	M	DW	100ML	3% H ₂ O ₂	539.6g	579.2g	39.6
3	GS	DRY	—	3% H ₂ O ₂	522.3g	535.7g	12.4
4	M	ADSORBENT**	~200GM.		678.9g	689.5g	10.6
#4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED			93.2
WEIGHT WITH ADSORBENT = _____ GM.							

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.67 ΔH_0 (POST-TEST) _____
Y (PRE-TEST) 1.02 Y (POST-TEST) _____

CONSOLE NO.
D- 328

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☐ THERMOCOUPLE (TYPE "K") ☐ OTHER _____	☐ STANDARD ☒ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER _____	☐ PYROMETER (TYPE "K") ☐ OTHER _____	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER <u>manometer</u>

COMMENTS:

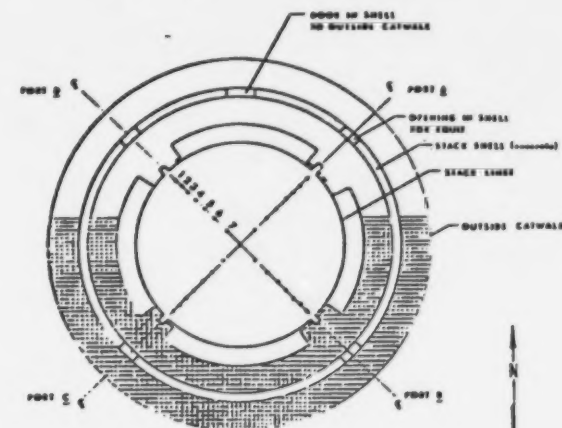
RUN NO. BEC-STK-7
DATE 7-22-80

KENVIRONS, INC.
STACK GEOMETRY

RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
(ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)

NOTE: SAME AS RUN _____



CATWALK INSIDE - OUTSIDE SHELL
10'-0" IS 11'00" AT PORT ELEV.
TOP OF BREACHING ELEV. 497'

PORT ELEV - 515'
PORT LENGTH - 10'
S.D. PORT W / CAP

DISTANCE FROM OUTSIDE PORT TO POINT (IN.)			
1. 10 1/2	2. 33 1/16	3. 93 13/16	4. 93 7/16
2. 33	4. 43 5/16	5. 91 5/16	

STACK
CROSS-SECTION

FOR ROUND STACKS

- A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT 238 IN.
B. PORT LENGTH 10 IN.
STACK DIAMETER: (A-B) 228 IN.
STACK AREA: 40828.1 IN.²

FOR RECTANGULAR STACKS

- A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT _____ IN.
B. PORT LENGTH _____ IN.
STACK DEPTH: (A-B) _____ IN.
STACK WIDTH (INSIDE) _____ IN.
STACK AREA (DEPTH X WIDTH) _____ IN.²
EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN.

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
DIAMETERS

11 DIAMETERS DOWNSTREAM OF
13 DIAMETERS UPSTREAM OF

TYPE OF FLOW
OBSTRUCTION

Breaching
top of stack

NUMBER OF
SAMPLING POINTS

MIN. REQ'D. (METHOD 1) 12
USE 28

RUN NO. REC-574-7
DATE 7/21/80

KENVIRONS, INC.
NOMOGRAPH DATA

RECORDED BY GPR
CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 87 °F
BAROMETRIC PRESSURE $P_b = 29.80$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 30.0) = 29.58$ IN. H₂O
 $\Delta H \theta$ REFERENCE ΔH = 1.60 IN. H₂O
TM AVG. METER TEMPERATURE = 107 °F
% H₂O MOISTURE CONTENT = 15 % BY VOL
PO STATIC (GUAGE) PRESSURE IN STACK = -0.75 IN. H₂O
x 0.074 = 0.06 IN. Hg.
PS ABSOLUTE PRESSURE IN STACK, $P_b - P_o$ = 29.52 IN. Hg.
PM METER PRESSURE ($\sim P_b$) = 29.56 IN. Hg.
PS/PM = _____
C CORRECTION FACTOR = _____
TS STACK TEMPERATURE = 175 °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O
D PROBE TIP DIAMETER = 0.25 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	100	125	150	175	
ΔP_{REF}					

PITOT LEAK CHECKS*				
OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	<u>GPR</u>	☒ STABLE ☐ LEAKING	<u>GPR</u>
STATIC	☒ STABLE ☐ LEAKING	<u>GPR</u>	☒ STABLE ☐ LEAKING	<u>GPR</u>

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS		
LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	<u>0.007 cfm @ 15" Hg</u>	<u>0.002 cfm @ 5.8" Hg</u>
TIME (24 HOUR CLOCK)	<u>10:33 thru tip</u>	<u>18:20 thru tip</u>
BY	<u>GPR + PM</u>	<u>GPR + PM</u>

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

ASSISTED BY Phm

SAMPLING TIME PER POINT 3 MIN.

SAMPLING TIME PER POINT 3 MIN.

E-255

KENVIRONS, INC.

RUN NO. BEC-STK-7 CARRIER GAS LOT
 DATE 7-23-80 SAMPLING (ORSAT) ASSISTED BY PM

NOTE: SAME AS RUN _____

A. GRAB SAMPLE (EVACUATED BULB)

SAMPLING TRAIN DESCRIPTION:

BULB MATERIAL AND VOLUME _____ ML.

PROBE MATERIAL AND LENGTH _____ IN.

PREFILTER MATERIAL _____

SAMPLING LOCATION: PORT _____ POINT _____

BULB NUMBER	INITIAL BULB VACUUM (IN. Hg)	VACUUM MEASUREMENT TIME (MIN.)	CLOCK TIME (24 HR. CLOCK)

B. INTEGRATED SAMPLE (TEDLAR BAG-TRAIN)

SAMPLING TRAIN DESCRIPTION:

PROBE MATERIAL AND LENGTH stainless steel 8 ft. IN.

PREFILTER MATERIAL none

PUMP TYPE diaphragm COOLING COIL stainless steel 6

SAMPLING LOCATION: PORT All POINT 2-7

SAMPLING TIME: START Same as method STOP _____

INTEGRATED SAMPLE TRAIN LEAK CHECKS*					
PRE-TEST	TIME	BY	POST-TEST	TIME	BY
☒ STABLE ☐ LEAKING	<u>16:05</u>	<u>PM</u>	☒ STABLE ☐ LEAKING	<u>1820</u>	<u>PM</u>

* Vacuum must remain stable for 30 seconds @ 10 in. Hg.

BAG LEAK CHECK PERFORMED BY RST TIME overnight
7/22/80 - 7/23/80

COMMENTS: _____

KENVIRONS, INC.
DRÄGER DATA FORM

RUN NO. BEC-STK-7
DATE 7-20-80

TYPE OF DRÄGER _____

RECORDED BY GR

ASSISTED BY _____

DISTANCE FROM DUCT INSIDE WALL TO DRÄGER SAMPLE TUBE TIP - ~12 IN.

DISTANCE FROM DUCT INSIDE WALL TO METHOD 6 OR 8 PROBE TIP - _____ IN.

SAMPLE PORT DESIGNATION C

ARE BOTH SAMPLES TAKEN FROM SAME PORT? YES NO

IF NO, APPROXIMATE DISTANCE FROM PROBE TIP TO SAMPLE TIP _____ IN.

WAS DRÄGER SAMPLE TUBE PURGED? ☒ YES NO

IF NO, GIVE REASON FOR NOT PURGING TUBE _____

RUN NO.	TIME	N NO. OF PUMPS FOR SAMPLE	R DRÄGER TUBE READING	R X 10 N IN PPM
STK-7A	18:40	10	160	160
STK-7B	18:45	10	150	150
Cal-A	18:50	10	90	90
Cal-B	18:55	10	120	120

} 101 ppm
SO₂
tank

COMMENTS: _____

RECORDED BY DM
ASSISTED BY _____

WAS FYRITE SAMPLE TUBE PURGED BEFORE SAMPLING? ☒ YES ☐ NO

IF NO, WHAT WAS THE REASON?

WAS FYRITE ZEROED INITIALLY? ✓ YES NO

WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ☒ YES ☐ NO

WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ☒ YES ☐ NO

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 12 IN.

[illegible]

COMMENTS:

RUN NO. BEC-STA-7 KENVIRONS, INC.
 DATE 7-23-80 CARRIER GAS
ANALYSIS (ORSAT) ANALYZED BY PR

ORSAT LEAK CHECKS											
PRE-ANALYSIS						POST-ANALYSIS					
NO.	TIME	BURETTE LEVEL		OK	BY	NO.	TIME	BURETTE LEVEL		OK	BY
	START	STOP	START				STOP	START	STOP		
1	1925 1930	17.3	17.3	✓	PR	1	2025 2030	35.6	35.6	✓	PR
2						2					
3						3					
4						4					

SAMPLE IDENTIFICATION	CO ₂		O ₂	READING 3	CO (3-2)
	READING 1	READING 2			
Stack #6	10.5	18.3	7.8	18.5	0.2
	10.5	18.4	7.9	18.5	0.1
	10.5	18.4	7.9	18.5	0.1
	AVG = 10.5		AVG = 7.867		AVG = 0.13
Stack #6	10.5	18.4	7.9	18.5	0.1
	10.6	18.4	7.8	18.5	0.1
	10.6	18.4	7.8	18.5	0.1
	AVG = 10.567		AVG = 7.833		AVG = 0.1
Ave.	10.53		7.95		0.12
	AVG =		AVG =		AVG =
	AVG =		AVG =		AVG =

CR
RST

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80014
FACILITY LGHE Can Run #6
RUN NUMBER BEC-57K-8

FIELD DATES 7-17-80/7-26-80
SAMPLING LOCATION Stack
SAMPLING DATE 7/24/80

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr.</u>
TESTING ENGINEER	1 <u>Gary H. Resett</u> 2 _____
ENGR. TECHNICIAN	1 <u>Parker Moore</u> 2 _____
LAB TECHNICIAN	1 _____ 2 _____
OTHER	1 _____ 2 _____

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-STK-4
DATE: 7/24/80

RECORDED BY: G/H
ASSISTED BY: _____

AMBIENT TEMPERATURE: _____ 80 _____ °F
BAROMETER PRESSURE: _____ 29.82 _____ in. Hg.
READING TIME: 7.55 RECORDED BY: RST
BAROMETER I.D.: 102 CALIBRATION DATE: 5/24/80
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-328
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: I
PROBE I.D.: 307 LENGTH: 8 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: VII IMPACT: side B STATIC: side A
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-3112
COLD BOX DIAL THERMOMETER I.D.: C - 1" Dial
HOT BOX DIAL THERMOMETER I.D.: C - 2" Dial
THERMOCOUPLE I.D.: 28 (Kearney Shield) LENGTH: 9 FT.
NOZZLE I.D.: 435 SIZE: 0.241
UMBILICAL I.D.: Green LENGTH: 25 FT.

COMMENTS: _____

RUN NO REC-STK-8
DATE 7-24-90

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY LR
ASSISTED BY _____

I. PROBE ASSEMBLY

A. PROBE LENGTH 8 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER VII
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.172 IN
B. NOZZLE IDENT. NO. 435
C. UMBILICAL LENGTH 75 F
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 79177

III. IMPINGER CONTENT

NUMBER	TYPE*	NORMAL		USE		FINAL AMOUNT	NET GAIN
		MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	520.8g	559.2g	38.4
2	M	DW	100ML	3% H ₂ O ₂	541.7g	587.7g	45.4
3	GS	DRY	_____	3% H ₂ O	524.6g	537.4g	13.3
4	M	ADSORBENT**	~200GM.		162.6g	764.6g	12.0
#4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED <u>109.1</u>			
WEIGHT WITH ADSORBENT = _____ GM.							

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.67 ΔH_0 (POST-TEST) _____

CONSOLE NO.

D- 328

Y (PRE-TEST) 1.02

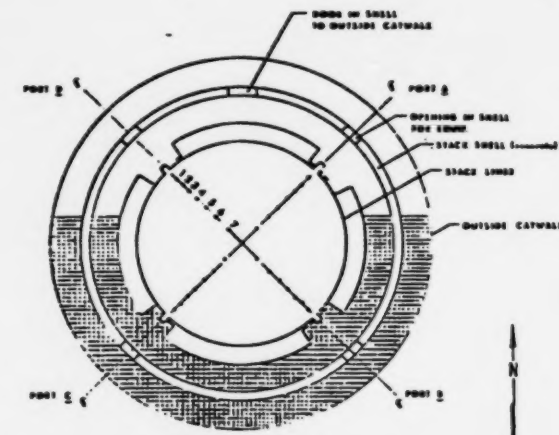
Y (POST-TEST) _____

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD	☐ THERMOCOUPLE (TYPE "K")	☐ STANDARD
	☒ "S" SHAPED	☐ OTHER _____	☒ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O)	☐ PYROMETER (TYPE "K")	☐ DI MANOMETER (0-5 IN. H ₂ O)
	☐ HIGH SENSITIVITY (RANGE _____)	☐ OTHER _____	☐ HIGH SENSITIVITY (RANGE _____)
	☐ OTHER _____		☐ OTHER <u>manometer</u>

COMMENTS: _____

RUN NO. BEC-STK-8 KENVIRONS, INC.
 DATE 2-24-80 STACK GEOMETRY RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
 (ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)
 NOTE: SAME AS RUN _____



CATWALL INSIDE - OUTSIDE SHELL
 10'-0" IN LINES AT PORT 1/2/3/4
 NEW 01 DISTANCE GUY. 400'
 PORT 1/2/3/4 - 710'
 PORT 1/2/3/4 - 10'
 A 0 PORT 1/2/3/4 CAP
 DISTANCE FROM OUTSIDE PORT TO POINT (IN)
 1. 00 5/16 2. 32 5/16 3. 00 10/16 4. 00 5/16
 5. 32 6. 40 5/16 7. 11 5/16

STACK
 CROSS-SECTION

FOR ROUND STACKS

A. DISTANCE FROM INSIDE FAR WALL
 TO OUTSIDE OF PORT 238 IN
 B. PORT LENGTH 10 IN
 STACK DIAMETER (A-B) 228 IN
 STACK AREA: 40828.1 IN²

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL
 TO OUTSIDE OF PORT _____ IN
 B. PORT LENGTH _____ IN
 STACK DEPTH (A-B) _____ IN
 STACK WIDTH (INSIDE) _____ IN
 STACK AREA (DEPTH X WIDTH) _____ IN²
 EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ _____ IN

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
 DIAMETERS

11 DIAMETERS DOWNSTREAM OF Breaching
13 DIAMETERS UPSTREAM OF top of stack

TYPE OF FLOW
 OBSTRUCTION

NUMBER OF
 SAMPLING POINTS

MIN. REQ'D. (METHOD 1) 12
 USE 28

RUN NO. BEC-STA-8
DATE 7/24/80

KENVIRONS, INC.
NOMOGRAPH DATA

RECORDED BY GHR
CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 80 °F

Pbc BAROMETRIC PRESSURE $P_b = \underline{29.82}$

ΔH CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 32.2) = \underline{29.60}$ IN. Hg.

TM REFERENCE $\Delta H = \underline{1.67}$ IN. H₂O

TM AVG. METER TEMPERATURE = 100 °F

% H₂O MOISTURE CONTENT = 15 % BY VOL

PO STATIC (GUAGE) PRESSURE IN STACK = -0.60 IN. H₂O

$\times 0.074 = \underline{0.05}$ IN. Hg.

PS ABSOLUTE PRESSURE IN STACK, $P_b - P_o = \underline{29.55}$ IN. Hg.

PM METER PRESSURE ($-P_b$) = 29.60 IN. Hg.

PS/PM = _____

C CORRECTION FACTOR = 0.77

TS STACK TEMPERATURE = 175 °F

ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O

ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O

D PROBE TIP DIAMETER = 0.24 / 0.175 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	100	125	150	175	
ΔP_{REF}				<u>29.82</u>	

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	<u>GHR</u>	☒ STABLE ☐ LEAKING	<u>GHR</u>
STATIC	☒ STABLE ☐ LEAKING	<u>GHR</u>	☒ STABLE ☐ LEAKING	<u>GHR</u>

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	<u>0.007 cfm @ 15" Hg</u>	<u>0.002 cfm @ 5" Hg</u>
TIME (24 HOUR CLOCK)	<u>10:50 hour</u>	<u>12:54 hour</u>
BY	<u>GHR</u>	<u>GHR</u>

* Rates < 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

KENVIRONS, INC.

RUN NO. BEC-STK-4

PARTICULATE

RECORDED BY GHRDATE 7/24/88

SAMPLING

ASSISTED BY PLSAMPLING TIME PER POINT 3 MIN.

PORT POINT	CLOCK TIME	METER VOLUME (DCF)	ΔP VEL. HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
A-1	11:00	101.52 108.41	0.45	0.26	92	84	162	275	68	3.4	
" 2		09.44	0.60	0.35	100	84	167	270	64	3.5	
" 3		10.55	0.70	0.41	105	84	168	265	62	3.9	
" 4		11.71	0.75	0.44	107	84	169	265	60	4.0	
" 5		12.88	0.75	0.44	107	84	169	260	59	4.1	
" 6		14.04	0.75	0.44	110	85	168	260	59	4.1	
A-7	11:21	15.20	0.75	0.44	111	85	168	265	62	4.1	
B-1	11:28	16.09	0.45	0.26	105	88	163	277	68	3.1	
" 2		17.12	0.60	0.35	110	88	169	270	66	3.7	
" 3		18.23	0.70	0.41	112	88	168	265	64	4.0	
" 4		19.40	0.75	0.44	113	88	167	270	68	4.2	
" 5		20.57	0.75	0.44	115	88	166	275	68	4.1	
" 6		21.74	0.75	0.44	115	89	166	280	66	4.2	
B-7	11:44	22.91	0.75	0.44	116	89	165	285	63	4.2	
C-1	11:55	23.74	0.45	0.26	105	91	167	285	68	3.2	
" 2		24.83	0.60	0.35	112	90	171	285	68	3.8	
" 3		25.97	0.70	0.41	115	90	174	280	75	4.0	
" 4		27.14	0.75	0.44	116	90	173	280	74	4.3	
" 5		28.33	0.75	0.44	116	90	169	275	78	4.3	
" 6		29.51	0.75	0.44	117	90	166	280	70	4.3	
C-7	12:16	30.70	0.75	0.44	118	90	167	280	67	4.3	

COMMENTS: 1.

9.

2.

10.

3.

11.

4.

12.

5.

13.

6.

14.

7.

15.

8.

16.

COMMENTS: 1.	9.
2.	10. <i>purged</i>
3.	11. <i>@ 0.40 ΔH</i>
4.	12.
5.	13.
6.	14.
7.	15.
8.	16.

RUN NO. BEC-STK-9 KENVIRONS, INC. CARRIER GAS 6R
 DATE 7-24-80 SAMPLING (ORSAT) ASSISTED BY _____

NOTE: SAME AS RUN _____

A. GRAB SAMPLE (EVACUATED BULB)

SAMPLING TRAIN DESCRIPTION:

BULB MATERIAL AND VOLUME _____ ML.

PROBE MATERIAL AND LENGTH _____ IN.

PREFILTER MATERIAL _____

SAMPLING LOCATION: PORT _____ POINT _____

BULB NUMBER	INITIAL BULB VACUUM (IN. Hg)	VACUUM MEASUREMENT TIME (MIN.)	CLOCK TIME (24 HR. CLOCK)

B. INTEGRATED SAMPLE (TEDLAR BAG-TRAIN)

SAMPLING TRAIN DESCRIPTION:

PROBE MATERIAL AND LENGTH stainless steel 8 ft. IN.

PREFILTER MATERIAL none

PUMP TYPE diaphragm COOLING COIL stainless steel in 6

SAMPLING LOCATION: PORT A11 POINT 2-7

SAMPLING TIME: START Same as method STOP 5

INTEGRATED SAMPLE TRAIN LEAK CHECKS*					
PRE-TEST	TIME	BY	POST-TEST	TIME	BY
☐ STABLE ☐ LEAKING	09:15	PM	☒ STABLE ☐ LEAKING	10:45	PM

* Vacuum must remain stable for 30 seconds @ 10 in. Hg.

BAG LEAK CHECK PERFORMED BY RST TIME overnight
7/23/80 - 7/24/80

COMMENTS: _____

KENVIRONS, INC.
DRÄGER DATA FORM

RUN NO. BEC-STK-8
DATE 7-24-80

TYPE OF DRÄGER _____

RECORDED BY GR
ASSISTED BY _____

DISTANCE FROM DUCT INSIDE WALL TO DRÄGER SAMPLE TUBE TIP - ~ 12 IN.

DISTANCE FROM DUCT INSIDE WALL TO METHOD 6 OR 8 PROBE TIP - — IN.

SAMPLE PORT DESIGNATION A and C

ARE BOTH SAMPLES TAKEN FROM SAME PORT? — YES — NO

IF NO, APPROXIMATE DISTANCE FROM PROBE TIP TO SAMPLE TIP — IN.

WAS DRÄGER SAMPLE TUBE PURGED? ✓ YES — NO

IF NO, GIVE REASON FOR NOT PURGING TUBE _____

RUN NO.	TIME	N NO. OF PUMPS FOR SAMPLE	R DRÄGER TUBE READING	R X 10 N IN PPM
STK-A	10:30	16	145	145
STK-B	13:10	16	125	125
STK-C	13:15	16	136	136

COMMENTS: _____

DATE 7/24/85

KENVIRONS, INC.
FYRITE DATA FORM

RECORDED BY P.H.M.

ASSISTED BY _____

WAS FYRITE SAMPLE TUBE PURGED BEFORE SAMPLING? ☒ YES ☐ NO

IF NO, WHAT WAS THE REASON?

WAS FYRITE ZEROED INITIALLY? ☒ YES ☐ NO

WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ☒ YES ☐ NO

WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ✓ YES NO

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 17 IN.

[illegible]

COMMENTS:

RUN NO. BEC-STA-8
 DATE 7-24-80

KENVIRONS, INC.
 CARRIER GAS
 ANALYSIS (ORSAT)

ANALYZED BY PRC

ORSAT LEAK CHECKS													
PRE-ANALYSIS						POST-ANALYSIS							
NO.	TIME		BURETTE LEVEL		OK	BY	NO.	TIME		BURETTE LEVEL		OK	BY
	START	STOP	START	STOP				START	STOP	START	STOP		
1	1502	1507	35.4	35.4	✓	PR	1	1552	1557	32.0	32.0	✓	PR
2							2						
3							3						
4							4						

SAMPLE IDENTIFICATION	CO ₂		O ₂	CO	
	READING 1	READING 2	(2-1)	READING 3	(3-2)
	10.8	18.4	7.6	18.5	.1
	10.9	18.5	7.7	18.5	0
	10.8	18.5	7.7	18.5	0
	AVG = 10.8		AVG = 7.7		AVG = 0
	10.8	18.6	7.6	18.6	0
	10.9	18.6	7.7	18.6	0
	10.9	18.6	7.7	18.6	0
	AVG = 10.9		AVG = 7.7		AVG = 0
Auc	10.85		7.7		0.0
	AVG =		AVG =		AVG =
	AVG =		AVG =		AVG =

correct
7.8

OK
RST

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80019
FACILITY LG+E Cam Run DC
RUN NUMBER BEC-STK-9

FIELD DATES 7-17-80 / 7-26-80
SAMPLING LOCATION Cam Run V6 Stack
SAMPLING DATE 7/25/80

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr.</u>
TESTING ENGINEER	<u>1 Gary Rowlett</u> <u>2</u>
ENGR. TECHNICIAN	<u>1 Parker Moore</u> <u>2</u>
LAB TECHNICIAN	<u>1</u> <u>2</u>
OTHER	<u>1</u> <u>2</u>

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BE-STK-8
DATE: 7/25/80

RECORDED BY: CHL
ASSISTED BY: _____

AMBIENT TEMPERATURE: _____ °F
BAROMETER PRESSURE: 29.84 _____ in. Hg.
READING TIME: 7:45 RECORDED BY: VAL
BAROMETER I.D.: 102 CALIBRATION DATE: 5/29/81
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-328
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: I
PROBE I.D.: 307 LENGTH: 8 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: VII IMPACT: side B STATIC: side A
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-3112
COLD BOX DIAL THERMOMETER I.D.: C - 1" Dial
HOT BOX DIAL THERMOMETER I.D.: C - 2" Dial
THERMOCOUPLE I.D.: 2B (Kenviros Shield) LENGTH: 9 FT.
NOZZLE I.D.: 435 SIZE: 0.241
UMBILICAL I.D.: Green LENGTH: 75 FT.

COMMENTS: _____

RUN NO REC-STK-9
DATE 7-25-80

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY 672
ASSISTED BY _____

I. PROBE ASSEMBLY

A. PROBE LENGTH 8 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER VII
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.192 IN
B. NOZZLE IDENT. NO. 435
C. UMBILICAL LENGTH 75 FT.
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER _____

III. IMPINGER CONTENT

NUMBER	TYPE*	NORMAL		USE		FINAL AMOUNT	NET GAIN
		MATERIAL	AMOUNT	MATERIAL	AMOUNT		
1	GS	DW	100ML	80% IPA	518.9g	554.8g	35.9g
2	M	DW	100ML	3% H ₂ O ₂	540.8g	586.8g	46.0g
3	GS	DRY	_____	3% H ₂ O	522.6g	531.7g	15.1g
4	M	ADSORBENT**	~200GM.		687.2g	700.2g	13.0g
#4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED _____			
WEIGHT WITH ADSORBENT = _____ GM.							

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.67 ΔH_0 (POST-TEST) _____
Y (PRE-TEST) 1.02 Y (POST-TEST) _____

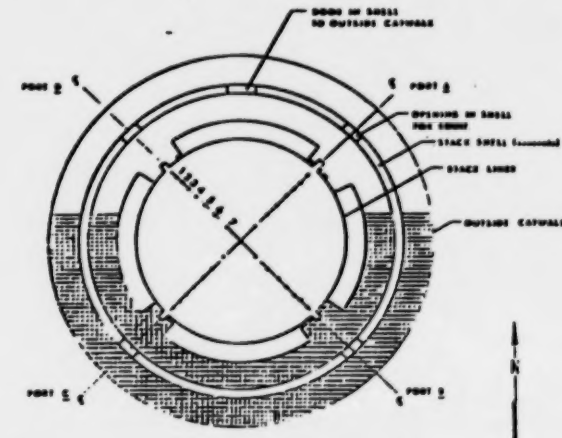
CONSOLE NO.
D- 328

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☐ THERMOCOUPLE (TYPE "K") ☐ OTHER _____	☐ STANDARD ☒ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER _____	☐ PYROMETER (TYPE "K") ☐ OTHER _____	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER <u>manometer</u>

COMMENTS: _____

RUN NO. BEC-STK-9 KENVIRONS, INC.
 DATE 7-25-80 STACK GEOMETRY RECORDED BY _____

SKETCH OF FLOW PROFILE AND CROSS-SECTION
 (ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)
 NOTE: SAME AS RUN _____



EATWALL INSIDE - OUTSIDE SHELL				PORT 1 (S1V - 75°)
IN - 10' 10" (10' 10" AT PORT 1 S1V)				PORT 1 (S1V - 10°)
TOP OF OUTSIDE SHELL 0.0'				0.0' PORT 1 W / E2P
DISTANCE FROM OUTSIDE PORT NO POINT (IN)				
1. 14 1/2"	3. 22 5/16"	5. 33 13/16"	7. 93 5/16"	
2. 23	4. 42 5/16"	6. 71 5/16"		

STACK
 CROSS-SECTION

FOR ROUND STACKS

- A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT 238 IN.
 B. PORT LENGTH 10 IN.
 STACK DIAMETER: (A-B) 228 IN.
 STACK AREA: 40828.1 IN²

FOR RECTANGULAR STACKS

- A. DISTANCE FROM INSIDE FAR WALL TO OUTSIDE OF PORT _____ IN.
 B. PORT LENGTH _____ IN.
 STACK DEPTH: (A-B) _____ IN.
 STACK WIDTH (INSIDE) _____ IN.
 STACK AREA (DEPTH X WIDTH) _____ IN.
 EQUIVALENT DIAMETER $\left(2 \sqrt{\frac{L \times W}{L + W}}\right)$ _____ IN.

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
 DIAMETERS

11 DIAMETERS DOWNSTREAM OF
13 DIAMETERS UPSTREAM OF

TYPE OF FLOW
 OBSTRUCTION

Breaching
top of stack

NUMBER OF
 SAMPLING POINTS

MIN. REQ'D. (METHOD 1) 12
 USE 28

RUN NO. BEC-STK-9 KENVIRONS, INC. RECORDED BY GHR
 DATE 7/25/80 NOMOGRAPH DATA CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 82 °F
 BAROMETRIC PRESSURE $P_b = 29.84$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 30) = 29.62$ IN. Hg
 ΔH REFERENCE $\Delta H = 1.60$ IN. H₂
 TM AVG. METER TEMPERATURE = 105 °F
 % H₂O MOISTURE CONTENT = 15 % BY
 P_0 STATIC (GUAGE) PRESSURE IN STACK = -0.60 IN. Hg
 x 0.074 = 0.06 IN. Hg
 PS ABSOLUTE PRESSURE IN STACK, $P_b - P_0 = 29.57$ IN. Hg
 PM METER PRESSURE ($\sim P_b$) = 29.62 IN. Hg
 PS/PM = _____
 C CORRECTION FACTOR = 0.775
 TS STACK TEMPERATURE = 175 °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂
 D PROBE TIP DIAMETER = 0.17 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	100	125	150	175	
ΔP_{REF}				3.1	

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	GHR	☒ STABLE ☐ LEAKING	GHR
STATIC	☒ STABLE ☐ LEAKING	GHR	☒ STABLE ☐ LEAKING	GHR

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	0.012 cfm @ 15 in. Hg	0.004 cfm @ 15 in. Hg
TIME (24 HOUR CLOCK)	10:25 thru tip	13:30 thru tip
BY	GHR - PM	GHR - PM

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

RECORDED BY Gofa

ASSISTED BY PA

SAMPLING TIME PER POINT 3 MIN.

PORT POINT	CLOCK TIME	METER VOLUME (DCF)	Δ P VEL. HEAD (IN. H ₂ O)	Δ H ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT
A - 1	11:45	44.33	0.50	0.29	90	85	173	250	64	3.4	
" 2		46.35	0.60	0.35	98	95	177	245	63	3.6	
" 3		47.52	0.75	0.44	104	96	173	230	61	4.0	
" 4		48.71	0.75	0.44	108	97	172	240	59	4.0	
" 5		49.92	0.80	0.47	110	98	171	245	59	4.2	
" 6		51.15	0.80	0.47	112	98	176	255	59	4.2	
A " 7	12:06	52.37	0.80	0.47	114	97	178	250	59	4.3	
D - 1	12:09	53.34	0.50	0.29	106	92	175	250	67	3.3	
" 2		54.43	0.65	0.38	114	92	179	245	66	3.8	
" 3		55.62	0.75	0.44	116	92	176	250	65	4.1	
" 4		56.84	0.80	0.47	118	92	171	245	63	4.2	
" 5		58.08	0.80	0.47	119	93	177	240	66	4.3	
" 6		59.30	0.80	0.47	120	94	172	250	62	4.3	
D " 7	12:30	60.54	0.80	0.47	121	94	171	250	64	4.3	
C - 1	12:35	61.50	0.50	0.29	110	96	174	250	68	3.4	
" 2		62.56	0.60	0.35	118	96	178	240	68	3.7	
" 3		63.73	0.80	0.47	120	96	181	250	67	4.3	
" 4		65.05	0.85	0.50	122	96	179	245	65	4.6	
" 5		66.32	0.85	0.50	122	96	178	240	65	4.6	
" 6		67.60	0.85	0.50	123	96	178	240	64	4.6	
C " 7	12:56	68.87	0.85	0.50	124	96	176	240	62	4.7	

COMMENTS: 1.

9.

2

10.

3

11.

4.

12.

5.

13.

6.

14.

7

15.

8.

16.

SAMPLING TIME PER POINT 3 MIN.

KENVIRONS, INC.

PARTICULATE

SAMPLING

RECORDED BY Gfr

ASSISTED BY Ph

[illegible]

COMMENTS: 1.

9.

purged 13:34-13

2

10.

⑥ $\Delta H \approx 0.45$

3.

11.

4.

12.

5.

13.

6.

14.

7.

15.

8.

16.

KENVIRONS, INC.
 RUN NO. BEC-STK-9 CARRIER GAS RECORDED BY GR
 DATE 7-25-80 SAMPLING (ORSAT) ASSISTED BY

NOTE: SAME AS RUN

A. GRAB SAMPLE (EVACUATED BULB)

SAMPLING TRAIN DESCRIPTION:

BULB MATERIAL AND VOLUME ML

PROBE MATERIAL AND LENGTH IN

PREFILTER MATERIAL

SAMPLING LOCATION: PORT POINT

BULB NUMBER	INITIAL BULB VACUUM (IN. Hg)	VACUUM MEASUREMENT TIME (MIN.)	CLOCK TIME (24 HR. CLOCK)

B. INTEGRATED SAMPLE (TEDLAR BAG-TRAIN)

SAMPLING TRAIN DESCRIPTION:

PROBE MATERIAL AND LENGTH stainless steel 8 ft. IN

PREFILTER MATERIAL none

PUMP TYPE diaphragm COOLING COIL stainless steel run

SAMPLING LOCATION: PORT A11 POINT 2-7

SAMPLING TIME: START same as method STOP

INTEGRATED SAMPLE TRAIN LEAK CHECKS*					
PRE-TEST	TIME	BY	POST-TEST	TIME	BY
☒ STABLE ☐ LEAKING	0:10:5	ph	☒ STABLE ☐ LEAKING	13:22	ph

* Vacuum must remain stable for 30 seconds @ 10 in. Hg.

BAG LEAK CHECK PERFORMED BY RS7 TIME overnight
7/24/80 - 7/25/80

COMMENTS:

RUN NO. BEC-STK-9
DATE - 7/25/80

RECORDED BY GHR
ASSISTED BY —

ARE BOTH SAMPLES TAKEN FROM SAME PORT? YES NO

IF NO, APPROXIMATE DISTANCE FROM PROBE TIP TO SAMPLE TIP _____ IN.
WAS DRÄGER SAMPLE TUBE PURGED? ☒ YES ☐ NO

IF NO, GIVE REASON FOR NOT PURGING TUBE

[illegible]

COMMENTS:

RECORDED BY JHM
ASSISTED BY _____

WAS FYRITE SAMPLE TUBE PURGED BEFORE SAMPLING? ☒ YES ☐ NO
IF NO, WHAT WAS THE REASON? _____

WAS FYRITE ZEROED INITIALLY? ☒ YES ☐ NO
 WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ☒ YES ☐ NO
 WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ☒ YES ☐ NO

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 12 IN.

[illegible]

COMMENTS: High O_2 in sample 1 caused by ambient air leaking into Fyrite tube thru filter connection. Leak repaired.

RUN NO. BEC-STX-
DATE 7-25-70

KENVIRONS, INC.
CARRIER GAS
ANALYSIS (ORSAT)

ANALYZED BY PR

ORSAT LEAK CHECKS											
PRE-ANALYSIS						POST-ANALYSIS					
NO.	TIME	BURETTE LEVEL		OK	BY	NO.	TIME	BURETTE LEVEL		OK	BY
	START STOP	START STOP	STOP				START STOP	START STOP	STOP		
1	1525 1530	32.0 32.0		✓	PR	1					
2						2					
3						3					
4						4					

SAMPLE IDENTIFICATION	CO ₂	READING 2	O ₂	READING 3	CO
	READING 1		(2-1)		(3-2)
Stack #6	10.7	18.5	7.8	18.6	—
	10.7	18.4	7.9	18.6	
	10.7	18.4	7.9	18.6	
	AVG = 10.7		AVG = 7.87		AVG = —
	10.7	18.4	7.9	18.6	—
	10.7	18.4	7.9	18.6	
	10.7	18.4	7.9	18.6	
	AVG = 10.7		AVG = 7.90		AVG = —
Aue	10.7		7.88		0
	AVG =		AVG =		AVG =
	AVG =		AVG =		AVG =

Ch
RST

BECHTEL ACCEPTANCE TEST

KENVIRONS, INC. PARTICULATE EMISSION TEST

PROJECT NUMBER 80014
FACILITY Bechtel
RUN NUMBER BEC-STK-10

FIELD DATES 7-17-80 / 7-26-80
SAMPLING LOCATION Cane Run #6 Stn
SAMPLING DATE 7/26/80

FIELD CREW

PROJECT CHIEF	<u>R.C. Tussey Jr.</u>
TESTING ENGINEER	1 <u>Gary H. Hewlett</u> 2 _____
ENGR. TECHNICIAN	1 <u>Parker Moore</u> 2 _____
LAB TECHNICIAN	1 _____ 2 _____
OTHER	1 _____ 2 _____

KENVIRONS, INC.
EQUIPMENT IDENTIFICATION

RUN NO.: BEC-STK-10
DATE: 7/26/86

RECORDED BY: Cif
ASSISTED BY: _____

AMBIENT TEMPERATURE: _____ °F
BAROMETER PRESSURE: 29.78 in. Hg.
READING TIME: 7:45 RECORDED BY: RST
BAROMETER I.D.: 107 CALIBRATION DATE: 5/29/86
CORRECTED BAROMETER PRESSURE: (@ console location)
CONSOLE I.D.: D-328
IN TEMP. DIAL THERMOMETER I.D.: II
OUT TEMP. DIAL THERMOMETER I.D.: I
PROBE I.D.: 307 LENGTH: 8 FT.
PROBE EXTENSION I.D.: _____ LENGTH: _____
PITOT I.D.: VII IMPACT: side B STATIC: side A
PITOT EXTENSION I.D.: _____ LENGTH: _____
SAMPLE BOX I.D.: S-3112
COLD BOX DIAL THERMOMETER I.D.: C - 1" Dial
HOT BOX DIAL THERMOMETER I.D.: C - 2" Dial
THERMOCOUPLE I.D.: 28 (Kenvirons Shield) LENGTH: 9 FT.
NOZZLE I.D.: 435 SIZE: 0.241
UMBILICAL I.D.: Green LENGTH: 75 FT.

COMMENTS: _____

RUN NO. BEC-STK-10
DATE 7-26-80

KENVIRONS, INC.
SAMPLING TRAIN
SPECIFICATIONS

RECORDED BY 612
ASSISTED BY _____

I. PROBE ASSEMBLY

A. PROBE LENGTH 8 FT.
B. LINER MATERIAL stainless steel
C. PITOT NUMBER VII
D. PITOT COEFFICIENT .84

II. MISC. COMPONENTS

A. NOZZLE DIAMETER 0.241
B. NOZZLE IDENT. NO. 435
C. UMBILICAL LENGTH 75
D. FILTER HOLDER TYPE 4" LSI
E. FILTER NUMBER 80042

III. IMPINGER CONTENT

		NORMAL		USE <i>642</i>		FINAL	NET
NUMBER	TYPE*	MATERIAL	AMOUNT	MATERIAL	AMOUNT	AMOUNT	GAIN
1	GS	DW	100ML	80% IPA	<i>518.2g</i>	530.9 g	12.7
2	M	DW	100ML	3% H ₂ O ₂	<i>536.9g</i>	587.8 g	50.9
3	GS	DRY	_____	3% H ₂ O	<i>526.5</i>	548.1 g	21.6
4	M	ADSORBENT**	~200GM.		<i>691.8g</i>	707.2 g	15.4
# 4 CONTAINER WEIGHT = _____ GM.				TOTAL MOISTURE ADSORBED		100.6	
WEIGHT WITH ADSORBENT = _____ GM.							

* GS = GREENBURG-SMITH, M = MODIFIED, O = OTHER

** SG = SILICA GEL, D = DRIERITE, O = OTHER

IV. CONTROL CONSOLE: ΔH_0 (PRE-TEST) 1.67 ΔH_0 (POST-TEST) _____
Y (PRE-TEST) 1.02 Y (POST-TEST) _____

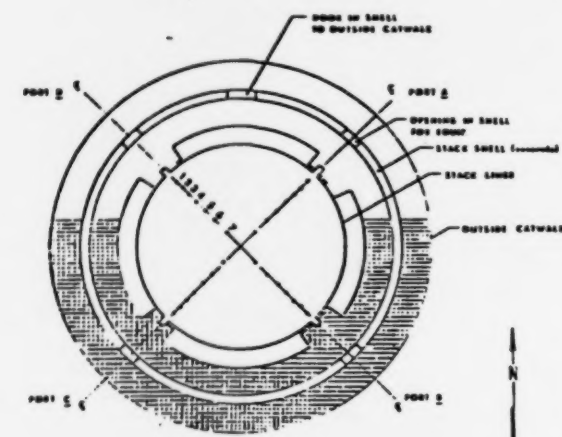
CONSOLE NO.
D- 328

	VELOCITY	TEMPERATURE	PRESSURE
SENSOR	☐ STANDARD ☒ "S" SHAPED	☐ THERMOCOUPLE (TYPE "K") ☐ OTHER _____	☐ STANDARD ☒ "S" SHAPED
READING DEVICE	☒ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER _____	☐ PYROMETER (TYPE "K") ☐ OTHER _____	☐ DI MANOMETER (0-5 IN. H ₂ O) ☐ HIGH SENSITIVITY (RANGE _____) ☐ OTHER <u>manometer</u>

COMMENTS: _____

RUN NO. BEC-STK-10 KENVIRONS, INC.
 DATE 7-26-80 STACK GEOMETRY RECORDED BY

SKETCH OF FLOW PROFILE AND CROSS-SECTION
 (ILLUSTRATE PORT-POINT CONFIGURATION, FLOW INTERFERENCES, ETC.)
 NOTE: SAME AS RUN



CATWALL INSIDE-OUTSIDE SHELL
 10'-0" IS LINES AT PORT 115°
 TOP OF BREACHING ELEV 402'

PORT 115°-710°
 PORT LENGTH-10"
 O.D. PORT W x CAP

DISTANCE FROM OUTSIDE PORT TO POINT (IN)			
1. 14 1/2"	2. 22 5/16"	3. 22 10/16"	4. 22 5/16"
5. 23"	6. 22 5/16"	7. 21 5/16"	

STACK
 CROSS-SECTION

FOR ROUND STACKS

A. DISTANCE FROM INSIDE FAR WALL
 TO OUTSIDE OF PORT 238 IN
 B. PORT LENGTH 10 IN
 STACK DIAMETER:(A-B) 228 IN
 STACK AREA: 40828.1 IN²

FOR RECTANGULAR STACKS

A. DISTANCE FROM INSIDE FAR WALL
 TO OUTSIDE OF PORT IN
 B. PORT LENGTH IN
 STACK DEPTH:(A-B) IN
 STACK WIDTH (INSIDE) IN
 STACK AREA(DEPTH X WIDTH) IN²
 EQUIVALENT DIAMETER $(2 \sqrt{\frac{L \times W}{L + W}})$ IN

SAMPLING CROSS-SECTION IS:

NO. OF DUCT
 DIAMETERS

11 DIAMETERS DOWNSTREAM OF
13 DIAMETERS UPSTREAM OF

TYPE OF FLOW
 OBSTRUCTION

Breaching
top of stack

NUMBER OF
 SAMPLING POINTS

MIN. REQ'D. (METHOD 1) 12
 USE 28

RUN NO. BEC-STX-10 KENVIRONS, INC. RECORDED BY GHA
 DATE 7/26/80 NOMOGRAPH DATA CHECKED BY _____

NOTE: SAME AS RUN _____

TA AMBIENT TEMPERATURE = 82 °F
 BAROMETRIC PRESSURE $P_b = 29.78$
 P_{bc} CORRECTED = $P_b - (.01 \text{ PER } 10 \text{ FT.}) \times (P_b - 30) = 29.56$ IN. Hg.
 ΔH REFERENCE $\Delta H = 1.60$ IN. H₂O
 TM AVG. METER TEMPERATURE = 102 °F
 $\% H_2O$ MOISTURE CONTENT = 15 % BY VOL
 P_0 STATIC (GUAGE) PRESSURE IN STACK = -0.75 IN. H₂O
 x 0.074 = 0.06 IN. Hg.
 P_S ABSOLUTE PRESSURE IN STACK, $P_b - P_0 = 29.50$ IN. Hg.
 P_M METER PRESSURE ($\sim P_b$) = 29.52 IN. Hg.
 $P_S/P_M =$ _____
 C CORRECTION FACTOR = 0.775
 T_S STACK TEMPERATURE = 170 °F
 ΔP AVERAGE STACK VELOCITY HEAD = _____ IN. H₂O
 ΔP_{MAX} MAXIMUM STACK VELOCITY HEAD = _____ IN. H₂O
 D PROBE TIP DIAMETER = 0.2416 IN.

Req'd Sampling Rate or Volume _____ Calculated Sampling Rate or Volume (Pre-test) _____

TS	100	125	150	175	
ΔP_{REF}					

PITOT LEAK CHECKS*

OPENING	PRE-TEST	BY	POST-TEST	BY
IMPACT	☒ STABLE ☐ LEAKING	<u>GHA</u>	☒ STABLE ☐ LEAKING	<u>GHA</u>
STATIC	☒ STABLE ☐ LEAKING	<u>GHA</u>	☒ STABLE ☐ LEAKING	<u>GHA</u>

* Vacuum must remain stable for 15 seconds @ 3 in. H₂O.

SAMPLING TRAIN LEAK CHECKS

LEAK CHECKS*	PRE-TEST	POST-TEST
RATE OF LEAK (CFM)	<u>0.017 cfm @ 15 in. Hg.</u>	<u>0.003 cfm @ 15 in. Hg.</u>
TIME (24 HOUR CLOCK)	<u>09:20 After Trp</u>	<u>11:24 After Trp</u>
BY	<u>LHR & PM</u>	<u>GHA & PM</u>

* Rates ≤ 0.02 CFM @ 15 in. Hg. vacuum or 4% of average sampling rate (whichever is less) are acceptable.

KENVIRONS, INC.
 RUN NO. REC-STK-10 PARTICULATE
 DATE 7/26/80 SAMPLING
 SAMPLING TIME PER POINT 3 MIN.
 RECORDED BY GHA
 ASSISTED BY PH

PORT POINT	CLOCK TIME	METER VOLUME (CCF)	ΔP VEL. HEAD (IN. H ₂ O)	ΔH ORIFICE (IN. H ₂ O)	METER TEMP IN (°F)	METER TEMP OUT (°F)	STACK TEMP (°F)	BOX TEMP (°F)	BATH TEMP (°F)	TRAIN VACUUM (IN. Hg)	COMMENT NUMBER
A-1	07:35	73.99	0.50	0.29	92	95	165	225	69	3.4	
" 2		84.09	0.65	0.38	103	85	180	240	67	3.5	
" 3		87.15	0.70	0.41	105	86	177	250	65	3.8	
" 4		88.31	0.75	0.44	109	86	175	260	67	4.0	
" 5		89.47	0.75	0.44	110	87	181	250	63	4.0	
" 6		90.64	0.75	0.44	113	88	183	255	64	4.0	
A-7	09:56	91.81	0.75	0.44	114	88	186	255	62	4.0	
D-1	10:01	92.76	0.80	0.29	107	92	169	270	70	3.2	
" 2		93.85	0.65	0.38	114	92	172	280	69	3.8	
" 3		94.94	0.70	0.41	116	92	177	280	69	3.9	
" 4		96.11	0.70	0.41	119	93	186	275	68	4.0	
" 5		97.29	0.75	0.44	120	94	192	275	68	4.0	
" 6		98.47	0.75	0.44	120	94	185	280	66	4.0	
D-7	10:22	99.65	0.75	0.44	121	94	178	275	67	4.0	
C-1	10:30	100.61	0.50	0.29	109	97	169	285	68	3.2	
" 2		101.71	0.65	0.38	118	96	171	295	68	3.8	
" 3		102.85	0.70	0.41	121	97	170	295	68	3.9	
" 4		104.00	0.70	0.41	122	97	175	295	67	4.0	
" 5		105.14	0.70	0.41	123	96	172	285	66	4.0	
" 6		106.29	0.70	0.41	124	97	170	290	66	4.0	
C-7	10:51	107.44	0.70	0.41	124	97	171	290	66	4.0	

COMMENTS-1.

2.

3.

4.

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16.

KENVIRONS, INC.
 RUN NO. BEC-STK-10 CARRIER GAS _____ RECORDED BY Gfr
 DATE 7/26/90 SAMPLING (ORSAT) _____ ASSISTED BY Ph

NOTE: SAME AS RUN _____

A. GRAB SAMPLE (EVACUATED BULB)

SAMPLING TRAIN DESCRIPTION:

BULB MATERIAL AND VOLUME _____ ML.

PROBE MATERIAL AND LENGTH _____ IN.

PREFILTER MATERIAL _____

SAMPLING LOCATION: PORT _____ POINT _____

BULB NUMBER	INITIAL BULB VACUUM (IN. Hg)	VACUUM MEASUREMENT TIME (MIN.)	CLOCK TIME (24 HR. CLOCK)

B. INTEGRATED SAMPLE (TEDLAR BAG-TRAIN)

SAMPLING TRAIN DESCRIPTION:

PROBE MATERIAL AND LENGTH stainless steel 8ft IN.

PREFILTER MATERIAL none

PUMP TYPE diaphragm COOLING COIL stainless steel sea 6

SAMPLING LOCATION: PORT All POINT 2-7

SAMPLING TIME: START same as method 5 STOP _____

INTEGRATED SAMPLE TRAIN LEAK CHECKS*					
PRE-TEST	TIME	BY	POST-TEST	TIME	BY
☒ STABLE ☐ LEAKING	09:00	PM	☐ STABLE ☐ LEAKING	11:20	PM

* Vacuum must remain stable for 30 seconds @ 10 in. Hg.

BAG LEAK CHECK PERFORMED BY RST/RR TIME overnight
7/25/90 - 7/26/90

COMMENTS: _____

RUN NO. BEC-STK-7D
DATE 7/26/90

TYPE OF DRAGER _____

RECORDED BY GHR
ASSISTED BY PM

ARE BOTH SAMPLES TAKEN FROM SAME PORT? YES NO

IF NO, APPROXIMATE DISTANCE FROM PROBE TIP TO SAMPLE TIP _____ IN.
WAS DRÄGER SAMPLE TUBE PURGED? ☒ YES ☐ NO

IF NO, GIVE REASON FOR NOT PURGING TUBE

[illegible]

COMMENTS: Only one Dräger taken, because only one was available.

Used 3 sets (10 ea.) during wk. @ 5th

(Apr. 1982) Rate 019/282

Dracopis CH 242, Cat. No. CH 24201 P¹

DATE 2/26/80

KENVIRONS, INC.
FYRITE DATA FORM

RECORDED BY PJM

ASSISTED BY _____

WAS FYRITE SAMPLE TUBE PURGED BEFORE SAMPLING? ☒ YES ☐ NO

IF NO, WHAT WAS THE REASON?

WAS FYRITE ZEROED INITIALLY? ☒ YES ☐ NO

WAS CO₂ FYRITE MIXED AT LEAST 4 TIMES? ☒ YES ☐ NO

WAS O₂ FYRITE MIXED AT LEAST 6 TIMES? ☒ YES ☐ NO

DISTANCE FROM INSIDE DUCT WALL TO END OF FYRITE SAMPLE TUBE 12 IN.

[illegible]

COMMENTS:

Analytical Particulate Filter Log

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REGISTRATION INFO (INITIAL)				TIME OF ARRIVAL ESTIMATION				REGISTRATION INFO (TIME)				FROM CLIMB ESTIMATION			
Serial Number	Flight Type	Project Number	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)	Time (10 min)
3173	79177		9:30	1:00	0946	VT	1205	1548	VT	1202					
3119	79178		9:30	1:02	0944	VT	1207	1545	VT	1202					
3117	79179		9:30	1:03	0947	VT	1208	1549	VT	1203					

5740-B
6870-A-B
6870-B-9

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REGISTRATION DATA (CONTINUED)										NAME REPORT INFORMATION										PHONE REPORT INFORMATION									
Source	Time	Lat	Long	Alt	Speed	Dir	Mag	Mag	Mag	Time	Lat	Long	Alt	Speed	Dir	Mag	Mag	Mag	Time	Lat	Long	Alt	Speed	Dir	Mag	Mag	Mag		
80023	4	80023	150	VT	11406	1019	VT	11402	1710	VT	11406	1019	VT	11402	1710	VT	11406	1019	VT	11406	1019	VT	11406	1019	VT	11406	1019	VT	
80024	"	"	"	"	11058	0951	"	11081	1142	"	11058	0951	"	11081	1142	"	11058	0951	"	11058	0951	"	11058	0951	"	11058	0951	"	
80025	"	"	"	"	11110	0955	"	11113	1146	"	11110	0955	"	11113	1146	"	11110	0955	"	11110	0955	"	11110	0955	"	11110	0955	"	
80026	"	"	"	"	11144	0956	"	11213	1145	"	11144	0956	"	11213	1145	"	11144	0956	"	11144	0956	"	11144	0956	"	11144	0956	"	
80027	"	"	"	"	11445	0957	"	11497	1149	"	11445	0957	"	11497	1149	"	11445	0957	"	11445	0957	"	11445	0957	"	11445	0957	"	
80028	"	"	"	"	11749	0958	"	1257	1148	"	11749	0958	"	1257	1148	"	11749	0958	"	11749	0958	"	11749	0958	"	11749	0958	"	
80029	"	"	"	"	11016	0959	"	11110	1148	"	11016	0959	"	11110	1148	"	11016	0959	"	11016	0959	"	11016	0959	"	11016	0959	"	
80030	"	"	"	"	11004	1000	"	11008	1150	"	11004	1000	"	11008	1150	"	11004	1000	"	11004	1000	"	11004	1000	"	11004	1000	"	
80031	"	"	"	"	11344	1001	"	11344	1152	"	11344	1001	"	11344	1152	"	11344	1001	"	11344	1001	"	11344	1001	"	11344	1001	"	
80032	"	"	"	"	11047	1002	"	11071	1151	"	11047	1002	"	11071	1151	"	11047	1002	"	11047	1002	"	11047	1002	"	11047	1002	"	
80033	"	"	"	"	11281	1003	"	11281	1153	"	11281	1003	"	11281	1153	"	11281	1003	"	11281	1003	"	11281	1003	"	11281	1003	"	
80034	"	"	"	"	11445	1004	"	11450	1154	"	11445	1004	"	11450	1154	"	11445	1004	"	11445	1004	"	11445	1004	"	11445	1004	"	
80035	"	"	"	"	11281	1005	"	11402	1155	"	11281	1005	"	11402	1155	"	11281	1005	"	11281	1005	"	11281	1005	"	11281	1005	"	
80036	"	"	"	"	11281	1006	"	11281	1156	"	11281	1006	"	11281	1156	"	11281	1006	"	11281	1006	"	11281	1006	"	11281	1006	"	
80037	"	"	"	"	11281	1008	"	11281	1157	"	11281	1008	"	11281	1157	"	11281	1008	"	11281	1008	"	11281	1008	"	11281	1008	"	
80038	"	"	"	"	11726	1020	"	11531	1109	"	11726	1020	"	11531	1109	"	11726	1020	"	11726	1020	"	11726	1020	"	11726	1020	"	
80039	"	"	"	"	11124	1044	"	11724	1138	"	11124	1044	"	11724	1138	"	11124	1044	"	11124	1044	"	11124	1044	"	11124	1044	"	
80040	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80041	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80042	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80043	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80044	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80045	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80046	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80047	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80048	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80049	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80050	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80051	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80052	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80053	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80054	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80055	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80056	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80057	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80058	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80059	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80060	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80061	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80062	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80063	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80064	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80065	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80066	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80067	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80068	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80069	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80070	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80071	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80072	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80073	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80074	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80075	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018	"	11064	1018	"	11064	1018	"	
80076	"	"	"	"	11064	1018	"	11077	1109	"	11064	1018	"	11077	1109	"	11064	1018	"	11064	1018								

REGISTRATION DATA (INITIAL)				TIME ELAPSED REGISTRATION				TIME ELAPSED REGISTRATION				TOTAL ELAPSED REGISTRATION				
Serial Number	Flight Type	Registration Number	Date	Time (Clock)	Time (Clock)	Time (Clock)	Time (Clock)	Date	Time (Clock)	Time (Clock)	Time (Clock)	Date	Time (Clock)	Time (Clock)	Time (Clock)	
3120	63			17-02-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87
3121	64			17-02-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87
3122	65			17-02-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87	1307	1207	1007	08-06-87

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Serial Number	Filter Type	REPRODUCTION DATA (100%)				TIME OF DAY REPRODUCTION				REPRODUCTION DATA (100%)				TIME OF DAY REPRODUCTION				REPRODUCTION DATA (100%)			
		Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)	Page	Time (min)
3013	1	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3014	2	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3015	3	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3016	4	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3017	5	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3018	6	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3019	7	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3020	8	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3021	9	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3022	10	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3023	11	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3024	12	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3025	13	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3026	14	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3027	15	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3028	16	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3029	17	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3030	18	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3031	19	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3032	20	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3033	21	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3034	22	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3035	23	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3036	24	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3037	25	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3038	26	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3039	27	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3040	28	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3041	29	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3042	30	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100
3043	31	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100

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KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Per-ESP-A-1 Date 7/29/80 By DAB/2nd (alt. 8/13/80) RM

=====

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3021 Final Beaker Weight: 102109.5 mg.
Acetone Volume: 109 ml. Beaker Tare Weight: 102109.4 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.1 mg. ÷ 109 ml. = 0.000917 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (785.7 mg/ml) × 100
% Blank Residual = 0.000917 mg/ml ÷ 785.7 mg/ml × 100 = 0.00012 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.000917 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No: Bec-ESP-A-1

ACETONE RINSE

Project No: 80019

Visual Observations: _____

By: DAB

Sample No.: 3024

Final Beaker Weight: 109740.1 mg.

Acetone Volume: 328 ml.

Beaker Tare Weight: 109332.5 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 407.6 mg.

Allowable Blank Residual

Minus Blank Correction: 0.3 mg.

Blank Correction =

Corrected Net Weight: 407.3 mg.

328 ml. x 0.00917 g/ml = 0.3 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations: _____

By: DAB

Filter No: 79149

Final Filter Weight: 1431.6 mg.

Sample No.: 3026

Filter Tare Weight: 1210.2 mg.

Filter Net Weight: 221.4 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 221.4 mg.

Sample Weight From Acetone Rinse: 407.3 mg.

Total Sample Weight: 628.7 mg.

Approved By: _____

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Per-ESPA-2 Date 7/31/80 By DAB/2nd Lab. 8-10-80 PAM

=====

ACETONE BLANK

Visual Observations: clear By: DAB
Sample No.: 3041 Final Beaker Weight: 100862.6 mg.
Acetone Volume: 100 ml. Beaker Tare Weight: 100861.9 mg.
Any Noted Acetone Loss: no Acetone Blank Net Weight: 0.7 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.7 mg. ÷ 100 ml. = 0.007 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (784.9 mg/ml) × 100
% Blank Residual = 0.007 mg/ml × 784.9 mg/ml × 100 = 0.00089 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.007 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = 0.00078 mg/ml

=====

=====

Run No. Bec-Esp-A-2 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3042 Final Beaker Weight: 109276.1 mg.

Acetone Volume: 330 ml. Beaker Tare Weight: 108838.0 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 438.1 mg.

Allowable Blank Residual Minus Blank Correction: 2.3 mg.

Blank Correction = Corrected Net Weight: 435.8 mg.

330 ml. x 0.007 mg/ml = 2.3 mg. Any Noted Acetone Loss: NONE

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79152 Final Filter Weight: 1623.0 mg.

Sample No.: 3040 Filter Tare Weight: 1236.7 mg.

Filter Net Weight: 386.3 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 386.3 mg.

Sample Weight From Acetone Rinse: 435.8 mg.

Total Sample Weight: 822.1 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

August 6, 1980

Analytical Log

Due to a malfunctioning valve in one of the desiccators the acetone evaporations from Day 2 may have been cross-contaminated, and some particulate lost in the desiccator. ESP-B-2 acetone rinse beaker was closest to the valve. Beakers ESP-A-2 acetone rinse and ESP-B-2 acetone blank were the only 2 beakers immediately adjacent to ESP-B-2 beaker. They received some loose dust from the ESP-B-2 spill. This loose dust, however, was deposited on top of firmly caked particulate matter in ESP-A-2 acetone rinse beaker and could be removed simply by pouring the loose dust back into ESP-B-2 acetone rinse beaker: loose dust also fell into ESP-B-2 acetone blank (empty) and was returned to ESP-B-2 acetone rinse in the same manner. The remainder of the loose dust was deposited within the desiccator itself and was returned to ESP-B-2 acetone rinse beaker using a camel hair brush and tweezers, although some particulate loss did occur. Below is a list of all beakers that were in the desiccator at the time of the spill, and the amount of contamination that each received:

Stk-2 Acetone Rinse	-	No Contamination
Stk-2 Acetone Blank	-	No Contamination
ESP-A-2 Acetone Rinse	-	Some Contamination
ESP-A-2 Acetone Blank	-	No Contamination
ESP-B-2 Acetone Rinse	-	Loss of Particulate
ESP-B-2 Acetone Blank	-	Some Contamination

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Ben-ESP-A-3 Date 7/31/80 By DAB/2nd Calc 8/13/80 PHL

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3050 Final Beaker Weight: 96943.1 mg.
Acetone Volume: 207 ml. Beaker Tare Weight: 96942.9 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.2 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.2 mg. + 207 ml. = 0.00097 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.7 mg/ml) x 100
% Blank Residual = 0.00097 mg/ml + 785.7 mg/ml x 100 = 0.00012 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00097 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = 0.0007857 mg/ml

=====

Run No. ESP-A-3

ACETONE RINSE

Project No. 80019

Visual Observations:

By: DAB

Sample No.: 3051

Final Beaker Weight: 102491.7 mg.

Acetone Volume: 440 ml.

Beaker Tare Weight: 101738.9 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 752.8 mg.

Allowable Blank Residual

Minus Blank Correction: 0.4 mg.

Blank Correction =

Corrected Net Weight: 752.4 mg.

440 ml. x 0.00097 mg/ml = 0.4 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 79156

Final Filter Weight: 1587.7 mg.

Sample No.: 3044

Filter Tare Weight: 1145.4 mg.

Filter Net Weight: 442.3 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 442.3 mg.

Sample Weight From Acetone Rinse: 752.4 mg.

Total Sample Weight: 1194.7 mg.

Approved By:

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Ben-ESP-A-4 Date 7/31/80 By DAB/2nd (24 8/13/80) JHM

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3061 Final Beaker Weight: 101576.4 mg.
Acetone Volume: 100 ml. Beaker Tare Weight: 101575.7 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.7 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.7 mg. ÷ 100 ml. = 0.007 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (7849 mg/ml) × 100
% Blank Residual = 0.007 mg/ml × 7849 mg/ml × 100 = 0.00089 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.007 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

Run No.: Bec-ESP-A-4

ACETONE RINSE

Project No.: 80019

Visual Observations:

By: DAB

Sample No.: 3065

Final Beaker Weight: 102335.7 mg.

Acetone Volume: 405 ml.

Beaker Tare Weight: 101610.1 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 725.6 mg.

Allowable Blank Residual

Minus Blank Correction: 2.8 mg.

Blank Correction =

Corrected Net Weight: 722.8 mg.

405 ml. x 0.007 mg/ml = 2.8 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No: 79159

Final Filter Weight: 15799 mg.

Sample No.: 3064

Filter Tare Weight: 1221.8 mg.

Filter Net Weight: 358.1 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 358.1 mg.

Sample Weight From Acetone Rinse: 722.8 mg.

Total Sample Weight: 1080.9 mg.

Approved By:

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bar-ESP-A-5 Date 7/31/80 By DAB/2nd Lab. 8:14:30 PM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3087 Final Beaker Weight: 108965.9 mg.
Acetone Volume: 100 ml. Beaker Tare Weight: 108965.8 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.1 mg. + 100 ml. = 0.001 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.5 mg/ml) x 100
% Blank Residual = 0.001 mg/ml + 785.5 mg/ml x 100 = 0.00013 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.001 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No.: Bec-ESP-A-5

ACETONE RINSE

Project No.: 80019

Visual Observations: _____

By: DAB

Sample No.: 3086

Final Beaker Weight: 100296.1 mg.

Acetone Volume: 370 ml.

Beaker Tare Weight: 99331.0 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 965.1 mg.

Allowable Blank Residual

Minus Blank Correction: 0.4 mg.

Blank Correction =

Corrected Net Weight: 964.7 mg.

370 ml. x 0.001 mg/ml = 0.4 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations: _____

By: DAB

Filter No.: 79168

Final Filter Weight: 1591.9 mg.

Sample No.: 3085

Filter Tare Weight: 1159.5 mg.

Filter Net Weight: 432.4 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 432.4 mg.

Sample Weight From Acetone Rinse: 964.7 mg.

Total Sample Weight: 1397.1 mg.

Approved By: _____

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESP-A-G Date 7/31/80 By DAB/2nd Calc. 8-14-80 PHM

ACETONE BLANK

Visual Observations:

By: DAB

Sample No.: 3097 Final Beaker Weight: 110765.5 mg.
Acetone Volume: 101 ml. Beaker Tare Weight: 110765.2 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.3 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume

Blank Residual = 0.3 mg. ÷ 101 ml. = 0.00297 mg/ml

% Blank Residual = Blank Residual ÷ Density of Acetone (7849 mg/ml) × 100

% Blank Residual = 0.00297 mg/ml ÷ 7849 mg/ml × 100 = 0.00038 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00297 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

Run No.: Bec-ESP-A-6

ACETONE RINSE

Project No.: 80019

Visual Observations:

By: DAB

Sample No.: 3096

Final Beaker Weight: 10879.6 mg.

Acetone Volume: 430 ml.

Beaker Tare Weight: 10795.3 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 834.3 mg.

Allowable Blank Residual

Minus Blank Correction: 1.3 mg.

Blank Correction =

Corrected Net Weight: 833.0 mg.

430 ml. x 0.003 mg/ml = 1.3 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 79172

Final Filter Weight: 1528.1 mg.

Sample No.: 3090

Filter Tare Weight: 1148.9 mg.

Filter Net Weight: 379.2 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 379.2 mg.

Sample Weight From Acetone Rinse: 833.0 mg.

Total Sample Weight: 1212.2 mg.

Approved By:

Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Be-ESP-A-7 Date 7/31/80 By DAB/2nd Lab. 9/13/80 RMM

ACETONE BLANK

Visual Observations: By: DAB
Sample No.: 3109 Final Beaker Weight: 99012.2 mg.
Acetone Volume: 116 ml. Beaker Tare Weight: 99011.8 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.4 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.4 mg. + 116 ml. = 0.00345 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.00345 mg/ml + 784.9 mg/ml x 100 = 0.00044 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00345 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

Run No: Ben-ESP 4-7 ACETONE RINSE Project No: 80019
 Visual Observations: _____ By: DAB
 Sample No.: 3110 Final Beaker Weight: 101380.1 mg.
 Acetone Volume: 430 ml. Beaker Tare Weight: 100909.0 mg.
 Blank Correction = Acetone Volume x Allowable Blank Residual
 Blank Correction = 430 ml. x 0.00345 mg/ml = 1.5 mg.
 Acetone Rinse Net Weight: 471.1 mg.
 Minus Blank Correction: 1.5 mg.
 Corrected Net Weight: 469.6 mg.
 Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations: _____ By: DAB
 Filter No: 79174 Final Filter Weight: 1442.4 mg.
 Sample No.: 3108 Filter Tare Weight: 11935 mg.
 Filter Net Weight: 2489 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 2489 mg.
 Sample Weight From Acetone Rinse: 469.6 mg.
 Total Sample Weight: 7185 mg.

Approved By: Gary H. Revlett Date: 8/14/80
 Gary H. Revlett, Ph.D.,
 Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESPA-8 Date 7/31/80 By DAB/2nd (alt. 9/13/80) PAM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3120 Final Beaker Weight: 97840.7 mg.
Acetone Volume: 98 ml. Beaker Tare Weight: 97840.2 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.5 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.5 mg. + 98 ml. = 0.0051 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.0051 mg/ml + 784.9 mg/ml x 100 = 0.00065 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.0051 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. ESP-A-8

ACETONE RINSE

Project No. 80019

Visual Observations:

By: DAB

Sample No.: 3124

Final Beaker Weight: 109970.4 mg.

Acetone Volume: 350 ml.

Beaker Tare Weight: 109462.9 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 5075 mg.

Allowable Blank Residual

Minus Blank Correction: 1.8 mg.

Blank Correction =

Corrected Net Weight: 505.7 mg.

350 ml. x 0.0051 mg/ml = 1.8 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 79178

Final Filter Weight: 1443.0 mg.

Sample No.: 3119

Filter Tare Weight: 1140.2 mg.

Filter Net Weight: 302.8 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 302.8 mg.

Sample Weight From Acetone Rinse: 505.7 mg.

Total Sample Weight: 808.5 mg.

Approved By:

Gary H. Revlett

Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-fsp-A-9 Date 7/31/80 By DAB/2nd Calc 8/13/80 JMM

ACETONE BLANK

Visual Observations: By: DAB
Sample No.: 3132 Final Beaker Weight: 100576.2 mg.
Acetone Volume: 64 ml. Beaker Tare Weight: 100575.9 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.3 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.3 mg. + 64 ml. = 0.00469 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.00469 mg/ml + 784.9 mg/ml x 100 = 0.000598%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00469 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

Run No. Bec-ESP-A-9 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3143 Final Beaker Weight: 109021.6 mg.

Acetone Volume: 470 ml. Beaker Tare Weight: 108435.6 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 586.0 mg.

Allowable Blank Residual Minus Blank Correction: 2.2 mg.

Blank Correction = Corrected Net Weight: 583.8 mg.

470 ml. x 0.00469 mg/ml = 2.2 mg. Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 80040 Final Filter Weight: 14240 mg.

Sample No.: 3134 Filter Tare Weight: 11004 mg.

Filter Net Weight: 323.6 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 323.6 mg.

Sample Weight From Acetone Rinse: 583.8 mg.

Total Sample Weight: 9074 mg.

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Esp-A-10 Date 7/31/80 By DAB

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3147 Final Beaker Weight: 104846.5 mg.
Acetone Volume: 102 ml. Beaker Tare Weight: 104846.4 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.1 mg. ÷ 102 ml. = 0.00098 mg/ml

% Blank Residual = Blank Residual ÷ Density of Acetone (784.9 mg/ml) × 100
% Blank Residual = 0.00098 mg/ml ÷ 784.9 mg/ml × 100 = 0.00012 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00098 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No. Bee-ESP-A-10

ACETONE RINSE

Project No. 80019

Visual Observations:

By: DAB

Sample No.: 3149

Final Beaker Weight: 102737.5 mg.

Acetone Volume: 540 ml.

Beaker Tare Weight: 101865.0 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 872.5 mg.

Allowable Blank Residual

Minus Blank Correction: 0.5 mg.

Blank Correction =

Corrected Net Weight: 872.0 mg.

540 ml. x 0.00098 mg/ml = 0.5 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 80043

Final Filter Weight: 1472.8 mg.

Sample No.: 3150

Filter Tare Weight: 1128.1 mg.

Filter Net Weight: 344.7 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 344.7 mg.

Sample Weight From Acetone Rinse: 872.0 mg.

Total Sample Weight: 1216.7 mg.

Approved By:

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

ESP-B-(1-10)

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESP-B-1 Date 7/29/80 By DAB/2nd Lab. 8/13-14/80 -TAM

ACETONE BLANK

Visual Observations:

By: DAB

Sample No.: 3022 Final Beaker Weight: 108966.3 mg.
Acetone Volume: 190 ml. Beaker Tare Weight: 108966.1 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.2 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume

Blank Residual = 0.2 mg. + 190 ml. = 0.00105 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.7 mg/ml) x 100

% Blank Residual = 0.00105 mg/ml + 785.7 mg/ml x 100 = 0.00013 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00105 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = mg/ml

Run No. Bec-ESP-B-1

ACETONE RINSE

Project No. : 80019

Visual Observations:

By: DAB

Sample No.: 3025

Final Beaker Weight: 101399.3 mg.

Acetone Volume: 443 ml.

Beaker Tare Weight: 100700.3 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 699.0 mg.

Allowable Blank Residual

Minus Blank Correction: 0.5 mg.

Blank Correction =

Corrected Net Weight: 698.5 mg.

443 ml. x 0.00125 mg/ml = 0.5 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 79148

Final Filter Weight: 1386.9 mg.

Sample No.: 3027

Filter Tare Weight: 1122.5 mg.

Filter Net Weight: 264.4 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 264.4 mg.

Sample Weight From Acetone Rinse: 698.5 mg.

Total Sample Weight: 962.9 mg.

Approved By: Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bac-Esp-B-2 Date 7/31/80 By DAB/2nd Lab 8-14-80 PAM

=====

ACETONE BLANK

Visual Observations: By: DAB
Sample No.: 3031 Final Beaker Weight: 101854.6 mg.
Acetone Volume: 180 ml. Beaker Tare Weight: 101854.0 mg.
Any Noted Acetone Loss: no Acetone Blank Net Weight: 0.6 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.6 mg. + 180 ml. = 0.00333 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (7849 mg/ml) x 100
% Blank Residual = 0.00333 mg/ml + 7849 mg/ml x 100 = 0.00042 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00333 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = 0.7849 mg/ml

=====

=====

Run No. Beo-Esp-B-2 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3038 Final Beaker Weight: 101163.5 mg.

Acetone Volume: 350 ml. Beaker Tare Weight: 100828.2 mg.

Blank Correction = Acetone Volume x _____ Acetone Rinse Net Weight: 335.3 mg.

Allowable Blank Residual _____ Minus Blank Correction: 1.2 mg.

Blank Correction = _____ Corrected Net Weight: 334.1 mg.

350 ml. x 0.00333 mg/ml = 1.2 mg. Any Noted Acetone Loss: NONE

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79150 Final Filter Weight: 1257.1 mg.

Sample No.: 3029 Filter Tare Weight: 1108.7 mg.

Filter Net Weight: 148.4 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 148.4 mg.

Sample Weight From Acetone Rinse: 334.1 mg.

Total Sample Weight: 482.5 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

August 6, 1980

Analytical Log

Due to a malfunctioning valve in one of the desiccators the acetone evaporations from Day 2 may have been cross-contaminated, and some particulate lost in the desiccator. ESP-B-2 acetone rinse beaker was closest to the valve. Beakers ESP-A-2 acetone rinse and ESP-B-2 acetone blank were the only 2 beakers immediately adjacent to ESP-B-2 beaker. They received some loose dust from the ESP-B-2 spill. This loose dust, however, was deposited on top of firmly caked particulate matter in ESP-A-2 acetone rinse beaker and could be removed simply by pouring the loose dust back into ESP-B-2 acetone rinse beaker: loose dust also fell into ESP-B-2 acetone blank (empty) and was returned to ESP-B-2 acetone rinse in the same manner. The remainder of the loose dust was deposited within the desiccator itself and was returned to ESP-B-2 acetone rinse beaker using a camel hair brush and tweezers, although some particulate loss did occur. Below is a list of all beakers that were in the desiccator at the time of the spill, and the amount of contamination that each received:

Stk-2 Acetone Rinse	-	No Contamination
Stk-2 Acetone Blank	-	No Contamination
ESP-A-2 Acetone Rinse	-	Some Contamination
ESP-A-2 Acetone Blank	-	No Contamination
ESP-B-2 Acetone Rinse	-	Loss of Particulate
ESP-B-2 Acetone Blank	-	Some Contamination

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Ber-ESP-B-3 Date 7/31/80 By DAB/2nd Calc. 8/13/80 PAM

=====

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3049 Final Beaker Weight: 104807.8 mg.
Acetone Volume: 187 ml. Beaker Tare Weight: 104807.6 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.2 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.2 mg. ÷ 187 ml. = 0.00107 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (785.7 mg/ml) × 100
% Blank Residual = 0.00107 mg/ml ÷ 785.7 mg/ml × 100 = 0.00014 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00107 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No. Bee-ESP-B-3 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3047 Final Beaker Weight: 10435.45 mg.

Acetone Volume: 470 ml. Beaker Tare Weight: 109235.8 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 1118.7 mg.

Allowable Blank Residual Minus Blank Correction: 0.5 mg.

Blank Correction = _____ Corrected Net Weight: 1118.2 mg.

470 ml. x 0.0007 mg/ml = 0.5 mg. Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 79154 Final Filter Weight: 1572.7 mg.

Sample No.: 3045 Filter Tare Weight: 1175.0 mg.

Filter Net Weight: 397.7 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 397.7 mg.

Sample Weight From Acetone Rinse: 1118.2 mg.

Total Sample Weight: 1515.9 mg.

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. FSP-B-4 Date 7/31/80 By DAB/2nd Lab. 8/13/80 PAM

=====

ACETONE BLANK

Visual Observations:

By: DAB

Sample No.: 3073 Final Beaker Weight: 101854.7 mg.
Acetone Volume: 120 ml. Beaker Tare Weight: 101853.9 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.8 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume

Blank Residual = 0.8 mg. + 120 ml. = 0.00667 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (7849 mg/ml) x 100

% Blank Residual = 0.00667 mg/ml + 7849 mg/ml x 100 = 0.00085 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00667 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No.: Bec-ESP-B-4

ACETONE RINSE

Project No.: 80019

Visual Observations:

By: DAB

Sample No.: 3071 ①

Final Beaker Weight: 98431.6 mg.

Acetone Volume: 450 ml.

Beaker Tare Weight: 97820.7 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 6109 mg.

Allowable Blank Residual

Minus Blank Correction: 3.0 mg.

Blank Correction =

Corrected Net Weight: 6079 mg.

450 ml. x 0.00667 mg/ml = 3.0 mg.

Any Noted Acetone Loss: none

① glass from broken funnel in sample

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No: 79167

Final Filter Weight: 1499.2 mg.

Sample No.: 3057

Filter Tare Weight: 1207.7 mg.

Filter Net Weight: 291.5 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 291.5 mg.

Sample Weight From Acetone Rinse: 6079 mg.

Total Sample Weight: 899.4 mg.

Approved By:

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESP-B-5 Date 7/31/80 By DAB / 2nd Calc. 9.11.80 FMM

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3075 Final Beaker Weight: 96942.7 mg.
Acetone Volume: 158 ml. Beaker Tare Weight: 96942.6 mg.
Any Noted Acetone Loss: _____ Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.1 mg. + 158 ml. = 0.0006 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.5 mg/ml) x 100
% Blank Residual = 0.0006 mg/ml + 785.5 mg/ml x 100 = 0.00008%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual 0.0006 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. Bec-ESP-B-5 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3074 Final Beaker Weight: 10271.7 mg.

Acetone Volume: 368 ml. Beaker Tare Weight: 10956.4 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 755.3 mg.

 Allowable Blank Residual Minus Blank Correction: 0.2 mg.

Blank Correction = Corrected Net Weight: 755.1 mg.

368 ml. x 0.0006 mg/ml = 0.2 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79169 Final Filter Weight: 1485.0 mg.

Sample No.: 3072 Filter Tare Weight: 1171.5 mg.

 Filter Net Weight: 313.5 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 313.5 mg.

Sample Weight From Acetone Rinse: 755.1 mg.

Total Sample Weight: 1068.6 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/96
Gary H. Revlett, PH.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Esp-B-6 Date 7/31/80 By DAB 2nd Calc. 8-14-80 PAM

=====

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3099 Final Beaker Weight: 103777.7 mg.
Acetone Volume: 355 ml. Beaker Tare Weight: 103777.2 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.5 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.5 mg. ÷ 355 ml. = 0.00141 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (7849 mg/ml) × 100
% Blank Residual = 0.00141 mg/ml ÷ 7849 mg/ml × 100 = 0.00018%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual
Allowable Blank Residual = 0.00141 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No. Bec-FSP B-6

ACETONE RINSE

Project No. 8009

Visual Observations: _____

By: DAB

Sample No.: 3098

Final Beaker Weight: 102742.2 mg.

Acetone Volume: 425 ml.

Beaker Tare Weight: 101452.9 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 1289.3 mg.

Allowable Blank Residual

Minus Blank Correction: 0.6 mg.

Blank Correction =

Corrected Net Weight: 1288.7 mg.

425 ml. x 0.0014 mg/ml = 0.6 mg.

Any Noted Acetone Loss: _____

FILTER WEIGHT

Visual Observations: _____

By: DAB

Filter No: 7170

Final Filter Weight: 1688.7 mg.

Sample No.: 3078

Filter Tare Weight: 1189.2 mg.

Filter Net Weight: 499.5 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 499.5 mg.

Sample Weight From Acetone Rinse: 1288.7 mg.

Total Sample Weight: 1788.2 mg.

Approved By: _____

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Esp-B-7 Date 7/31/80 By DAB/2nd Calc. 8/13/80 JHM

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3102 Final Beaker Weight: 110446.0 mg.
Acetone Volume: 130 ml. Beaker Tare Weight: 110445.1 mg.
Any Noted Acetone Loss: _____ Acetone Blank Net Weight: 0.9 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume

Blank Residual = 0.9 mg. ÷ 130 ml. = 0.0069 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (784.9 mg/ml) × 100
% Blank Residual = 0.0069 mg/ml ÷ 784.9 mg/ml × 100 = 0.00088 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.0069 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No. BeeESP-B-7 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3103 Final Beaker Weight: 101332.8 mg.

Acetone Volume: 400 ml. Beaker Tare Weight: 100700.1 mg.

Blank Correction = Acetone Volume x _____ Acetone Rinse Net Weight: 632.7 mg.

Allowable Blank Residual _____ Minus Blank Correction: 2.8 mg.

Blank Correction = _____ Corrected Net Weight: 629.9 mg.

400 ml. x 0.007 mg/ml = 2.8 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79175 Final Filter Weight: 1546.6 mg.

Sample No.: 3092 Filter Tare Weight: 1254.8 mg.

Filter Net Weight: 291.8 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 291.8 mg.

Sample Weight From Acetone Rinse: 629.9 mg.

Total Sample Weight: 921.7 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESP-B-8 Date 7/31/80 By DAB/2nd Calc. 8/13/80 RHM

=====

ACETONE BLANK

Visual Observations: By: DAB
Sample No.: 3122 Final Beaker Weight: 102145.2 mg.
Acetone Volume: 110 ml. Beaker Tare Weight: 102144.8 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.4 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.4 mg. + 110 ml. = 0.0036 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.0036 mg/ml + 784.9 mg/ml x 100 = 0.00046%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.0036 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. Ben ESP-B-8 ACETONE RINSE Project No. 8009

Visual Observations: _____ By: DAB

Sample No.: 3118 Final Beaker Weight: 97921.2 mg.

Acetone Volume: 375 ml. Beaker Tare Weight: 97400.9 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 520.3 mg.

Allowable Blank Residual Minus Blank Correction: 1.4 mg.

Blank Correction = Corrected Net Weight: 518.9 mg.

375 ml. x 0.0036 mg/ml = 1.4 mg. Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79176 Final Filter Weight: 1469.5 mg.

Sample No.: 3105 Filter Tare Weight: 1182.8 mg.

Filter Net Weight: 286.7 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 286.7 mg.

Sample Weight From Acetone Rinse: 518.9 mg.

Total Sample Weight: 805.6 mg.

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-ESP-B-9 Date 7/31/80 By DAB/2nd Calc 9/13/84 DAB

ACETONE BLANK

Visual Observations:

By: DAB

Sample No.: 3136 Final Beaker Weight: 104807.8 mg.

Acetone Volume: 226 ml. Beaker Tare Weight: 104807.7 mg.

Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume

Blank Residual = 0.1 mg. ÷ 226 ml. = 0.000442 mg/ml

% Blank Residual = Blank Residual ÷ Density of Acetone (784.9 mg/ml) × 100

% Blank Residual = 0.000442 mg/ml ÷ 784.9 mg/ml × 100 = 0.000056 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.000442 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

Run No. BEC-ESP-B9 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3137 Final Beaker Weight: 100576.6 mg.

Acetone Volume: 540 ml. Beaker Tare Weight: 99650.0 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 926.6 mg.

Allowable Blank Residual Minus Blank Correction: 0.2 mg.

Blank Correction = _____ Corrected Net Weight: 926.4 mg.

540 ml. x 0.00042 mg/ml = 0.2 mg. Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 79179 Final Filter Weight: 1459.1 mg.

Sample No.: 3117 Filter Tare Weight: 1143.3 mg.

Filter Net Weight: 315.8 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 315.8 mg.

Sample Weight From Acetone Rinse: 926.2 mg.

Total Sample Weight: 1242.0 mg.

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Rec-ESP-B-10 Date 7/31/80 By DAB

=====

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3145 Final Beaker Weight: 104248.2 mg.
Acetone Volume: 130 ml. Beaker Tare Weight: 104247.7 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.5 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.5 mg. + 130 ml. = 0.00385 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.00385 mg/ml + 784.9 mg/ml x 100 = 0.00049 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00385 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

=====

Run No. Bec-ESP-B-10 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3144 Final Beaker Weight: 109907.3 mg.

Acetone Volume: 465 ml. Beaker Tare Weight: 109414.2 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 493.1 mg.

Allowable Blank Residual Minus Blank Correction: 1.8 mg.

Blank Correction = 385 Corrected Net Weight: 491.3 mg.

465 ml. x 0.00385 mg/ml = 1.8 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DA13

Filter No: 80044 Final Filter Weight: 13634 mg.

Sample No.: 3142 Filter Tare Weight: 1144.5 mg.

Filter Net Weight: 218.9 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 218.9 mg.

Sample Weight From Acetone Rinse: 491.3 mg.

Total Sample Weight: 710.2 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/86
Gary H. Revlett, Ph.D.,
Analytical Chemist

STK-(1-10)

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bea Stk-1 Date 7/29/80 By DAB / 2nd Calc 8/13-14-7MM
OK DAB

ACETONE BLANK

Visual Observations: clear By: DAB
Sample No.: 3020 Final Beaker Weight: 108552.6 mg.
Acetone Volume: 208 ml. Beaker Tare Weight: 108552.4 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.2 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume

Blank Residual = 0.2 mg. + 208 ml. = 0.00096 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.7 mg/ml) x 100

% Blank Residual = 0.00096 mg/ml + 785.7 mg/ml x 100 = 0.00012 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00096 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = mg/ml

=====

Run No. Bec-Stk-1 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3023 Final Beaker Weight: 107167.9 mg.

Acetone Volume: 454 ml. Beaker Tare Weight: 107144.0 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 23.9 mg.

Allowable Blank Residual Minus Blank Correction: 0.4 mg.

Blank Correction = Corrected Net Weight: 23.5 mg.

454 ml. x 0.0009 mg/ml = 0.4 mg. Any Noted Acetone Loss: NONE

=====

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 79146 Final Filter Weight: 1213.6 mg.

Sample No.: 3028 Filter Tare Weight: 1157.3 mg.

Filter Net Weight: 56.3 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 56.3 mg.

Sample Weight From Acetone Rinse: 23.5 mg.

Total Sample Weight: 79.8 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/88
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-2 Date 7/29/80 By DAB/2nd Lab 3:14 PM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3043 Final Beaker Weight: 99309.5 mg.
Acetone Volume: 140 ml. Beaker Tare Weight: 99309.2 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.3 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.3 mg. ÷ 140 ml. = 0.002143 mg/ml

% Blank Residual = Blank Residual ÷ Density of Acetone (784.9 mg/ml) × 100
% Blank Residual = 0.002143 mg/ml ÷ 784.9 mg/ml × 100 = 0.00027%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.002143 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

=====

Run No.: BCC-STK-2

ACETONE RINSE

Project No.: 90019

Visual Observations:

By: DAB

Sample No.: 3037

Final Beaker Weight: 108104.4 mg.

Acetone Volume: 370 ml.

Beaker Tare Weight: 108069.4 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 350 mg.

Allowable Blank Residual

Minus Blank Correction: 0.8 mg.

Blank Correction =

Corrected Net Weight: 34.2 mg.

370 ml. x 0.0001142 mg/ml = 0.8 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No.: 79151

Final Filter Weight: 1291.1 mg.

Sample No.: 3039

Filter Tare Weight: 1232.6 mg.

Filter Net Weight: 58.5 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 58.5 mg.

Sample Weight From Acetone Rinse: 32.4 mg.

Total Sample Weight: 90.9 mg.

Approved By:

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/88

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 8009
Run No. Bec-Stk-3 Date 7/31/80 By DAB/2nd Calc. 8/13/80 AWM

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3057 Final Beaker Weight: 108672.6 mg.
Acetone Volume: 192 ml. Beaker Tare Weight: 108672.3 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.3 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume

Blank Residual = 0.3 mg. + 192 ml. = 0.00156 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.7 mg/ml) x 100

% Blank Residual = 0.00156 mg/ml + 785.7 mg/ml x 100 = 0.00020 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00156 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. BEC-SHK-3 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3058 Final Beaker Weight: 107517.4 mg.

Acetone Volume: 450 ml. Beaker Tare Weight: 107481.0 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 36.4 mg.

Allowable Blank Residual Minus Blank Correction: 0.7 mg.

Blank Correction = Corrected Net Weight: 35.7 mg.

450 ml. x 0.00156 mg/ml = 0.7 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No: 79153 Final Filter Weight: 1277.3 mg.

Sample No.: 3052 Filter Tare Weight: 1216.3 mg.

Filter Net Weight: 61.0 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 61.0 mg.

Sample Weight From Acetone Rinse: 35.7 mg.

Total Sample Weight: 96.7 mg.

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Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-4 Date 7/31/87 By DAB/2nd Calc. 9/13-14/80 R/M

ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3068 Final Beaker Weight: 111083.8 mg.
Acetone Volume: 185 ml. Beaker Tare Weight: 111082.9 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.9 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.9 mg. + 185 ml. = 0.004865 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.004865 mg/ml + 784.9 mg/ml x 100 = 0.00062 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.004865 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

Run No.: Bec-Stk-4

ACETONE RINSE

Project No.: 60019

Visual Observations: _____

By: DAB

Sample No.: 3069

Final Beaker Weight: 101957.6 mg.

Acetone Volume: 340 ml.

Beaker Tare Weight: 101923.3 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 34.3 mg.

Allowable Blank Residual

Minus Blank Correction: 1.7 mg.

Blank Correction =

Corrected Net Weight: 32.6 mg.

340 ml. x 0.00485 mg/ml = 1.7 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations: _____

By: DAB

Filter No.: 79158

Final Filter Weight: 1290.6 mg.

Sample No.: 3060

Filter Tare Weight: 1232.0 mg.

Filter Net Weight: 58.6 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 58.6 mg.

Sample Weight From Acetone Rinse: 32.6 mg.

Total Sample Weight: 91.2 mg.

Approved By: Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 9/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Ber-Stk-5 Date 7/31/80 By DAB/2nd Calc. 9-14-80 ZHM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3084 Final Beaker Weight: 102109.9 mg.
Acetone Volume: 204 ml. Beaker Tare Weight: 102109.5 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.4 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.4 mg. + 204 ml. = 0.00196 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (785.5 mg/ml) x 100
% Blank Residual = 0.00196 mg/ml + 785.5 mg/ml x 100 = 0.00025 %

If Blank Residual is less than 0.001% then:
Allowable Blank Residual = Blank Residual
Allowable Blank Residual = 0.00196 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:
Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

=====

Run No. Bee-Stk-5 ACETONE RINSE Project No. 8009

Visual Observations: _____ By: DAB

Sample No.: 3088 Final Beaker Weight: 109782.8 mg.

Acetone Volume: 300 ml. Beaker Tare Weight: 109752.2 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 30.6 mg.

Allowable Blank Residual Minus Blank Correction: 0.6 mg.

Blank Correction = Corrected Net Weight: 30.0 mg.

300 ml. x 0.0096 mg/ml = 0.6 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 79160 Final Filter Weight: 12570 mg.

Sample No.: 3077 Filter Tare Weight: 1196.7 mg.

Filter Net Weight: 60.3 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 60.3 mg.

Sample Weight From Acetone Rinse: 30.0 mg.

Total Sample Weight: 90.3 mg.

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Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-6 Date 7/31/80 By DAB/2nd Calc. 8-14-80 PHL

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ACETONE BLANK

Visual Observations: _____ By: DAB

Sample No.: 3100 Final Beaker Weight: 101272.9 mg.
Acetone Volume: 189 ml. Beaker Tare Weight: 101272.1 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.8 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume

Blank Residual = 0.8 mg. + 189 ml. = 0.0042 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100

% Blank Residual = 0.0042 mg/ml + 784.9 mg/ml x 100 = 0.00051%

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.0042 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. Bec-Stk-6

ACETONE RINSE

Project No. 80019

Visual Observations:

By: DAB

Sample No.: 3101

Final Beaker Weight: 100926.0 mg.

Acetone Volume: 325 ml.

Beaker Tare Weight: 100888.2 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 37.8 mg.

Allowable Blank Residual

Minus Blank Correction: 1.4 mg.

Blank Correction =

Corrected Net Weight: 36.4 mg.

325 ml. x 0.0042 mg/ml = 1.4 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No: 79171

Final Filter Weight: 1266.9 mg.

Sample No.: 3079

Filter Tare Weight: 1212.0 mg.

Filter Net Weight: 54.9 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 54.9 mg.

Sample Weight From Acetone Rinse: 36.4 mg.

Total Sample Weight: 91.3 mg.

Approved By: Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/80

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-7 Date 7/31/80 By DAB/2nd Calc. 8/13/80 PWH

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3114 Final Beaker Weight: 108553.8 mg.
Acetone Volume: 180 ml. Beaker Tare Weight: 108553.1 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.7 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.7 mg. + 180 ml. = 0.00389 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.00389 mg/ml + 784.9 mg/ml x 100 = 0.000495

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00389 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No.: Ber-Stk-7

ACETONE RINSE

Project No.: 86819

Visual Observations:

By: DAB

Sample No.: 3115

Final Beaker Weight: 101120.5 mg.

Acetone Volume: 375 ml.

Beaker Tare Weight: 101082.1 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 38.4 mg.

Allowable Blank Residual

Minus Blank Correction: 1.5 mg.

Blank Correction =

Corrected Net Weight: 36.9 mg.

375 ml. x 0.00389 mg/ml = 1.5 mg.

Any Noted Acetone Loss: NONE

FILTER WEIGHT

Visual Observations:

By: DAB

Filter No: 79173

Final Filter Weight: 1251.2 mg.

Sample No.: 3107

Filter Tare Weight: 1207.0 mg.

Filter Net Weight: 44.2 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 44.2 mg.

Sample Weight From Acetone Rinse: 36.9 mg.

Total Sample Weight: 81.1 mg.

Approved By:

Harry A. Revlett
Harry A. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/60

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-8 Date 7/31/80 By DAB / 2nd Calc. 8/13/80 PDM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3126 Final Beaker Weight: 996344 mg.
Acetone Volume: 186 ml. Beaker Tare Weight: 996343 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.1 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.1 mg. + 186 ml. = 0.00054 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.00054 mg/ml + 784.9 mg/ml x 100 = 0.00069 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.00054 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

Run No. Bec STK-8

ACETONE RINSE

Project No. 80019

Visual Observations: _____

By: DAB

Sample No.: 3125

Final Beaker Weight: 101057.6 mg.

Acetone Volume: 260 ml.

Beaker Tare Weight: 101028.4 mg.

Blank Correction = Acetone Volume x

Acetone Rinse Net Weight: 29.2 mg.

Allowable Blank Residual

Minus Blank Correction: 0.1 mg.

Blank Correction =

Corrected Net Weight: 29.1 mg.

260 ml. x 0.00054 mg/ml = 0.1 mg.

Any Noted Acetone Loss: none

FILTER WEIGHT

Visual Observations: _____

By: DAB

Filter No.: 79177

Final Filter Weight: 1185.2 mg.

Sample No.: 3123

Filter Tare Weight: 1138.0 mg.

Filter Net Weight: 47.2 mg.

SAMPLE WEIGHT

Sample Weight From Filter: 47.2 mg.

Sample Weight From Acetone Rinse: 29.1 mg.

Total Sample Weight: 76.3 mg.

Approved By: _____

Gary H. Revlett
Gary H. Revlett, Ph.D.,
Analytical Chemist

Date: 8/14/90

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-9 Date 7/31/80 By DAB/2nd Calc 8/13/80 PAM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3139 Final Beaker Weight: 97634.0 mg.
Acetone Volume: 94 ml. Beaker Tare Weight: 97633.3 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.7 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.7 mg. + 94 ml. = 0.007447 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.007447 mg/ml + 784.9 mg/ml x 100 = 0.00095 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.007447 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

=====

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Run No. Bec-Stk-9 ACETONE RINSE Project No. 80019

Visual Observations: _____ By: DAB

Sample No.: 3138 Final Beaker Weight: 106633.1 mg.

Acetone Volume: 292 ml. Beaker Tare Weight: 1066000 mg.

Blank Correction = Acetone Volume x Acetone Rinse Net Weight: 33.1 mg.

Allowable Blank Residual Minus Blank Correction: 2.2 mg.

Blank Correction = _____ Corrected Net Weight: 30.9 mg.

292 ml. x 0.0074 mg/ml = 2.2 mg. Any Noted Acetone Loss: none

=====

FILTER WEIGHT

Visual Observations: _____ By: DAB

Filter No.: 80041 Final Filter Weight: 1203.9 mg.

Sample No.: 3130 Filter Tare Weight: 1136.4 mg.

Filter Net Weight: 67.5 mg.

=====

SAMPLE WEIGHT

Sample Weight From Filter: 67.5 mg.

Sample Weight From Acetone Rinse: 30.9 mg.

Total Sample Weight: 98.4 mg.

=====

Approved By: Gary H. Revlett Date: 8/14/80
Gary H. Revlett, Ph.D.,
Analytical Chemist

KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Ben-Stk-10 Date 7/31/80 By DAB

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3154 Final Beaker Weight: 108672.2 mg.
Acetone Volume: 188 ml. Beaker Tare Weight: 108672.0 mg.
Any Noted Acetone Loss: NONE Acetone Blank Net Weight: 0.2 mg.

Blank Residual = Acetone Blank Net Weight ÷ Acetone Volume
Blank Residual = 0.2 mg. ÷ 188 ml. = 0.0011 mg/ml

% Blank Residual = Blank Residual × Density of Acetone (784.9 mg/ml) × 100
% Blank Residual = 0.0011 mg/ml ÷ 784.9 mg/ml × 100 = 0.00014 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.0011 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone × 0.00001 = _____ mg/ml

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KENVIRONS, INC.
EPA METHOD 5 LABORATORY
DATA SHEET

Client Bechtel Project No. 80019
Run No. Bec-Stk-9 Date 7/31/80 By DAB/2nd Calc 8/13/80 PAM

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ACETONE BLANK

Visual Observations: _____ By: DAB
Sample No.: 3139 Final Beaker Weight: 97634.0 mg.
Acetone Volume: 94 ml. Beaker Tare Weight: 97633.3 mg.
Any Noted Acetone Loss: none Acetone Blank Net Weight: 0.7 mg.

Blank Residual = Acetone Blank Net Weight + Acetone Volume
Blank Residual = 0.7 mg. + 94 ml. = 0.007447 mg/ml

% Blank Residual = Blank Residual + Density of Acetone (784.9 mg/ml) x 100
% Blank Residual = 0.007447 mg/ml + 784.9 mg/ml x 100 = 0.00095 %

If Blank Residual is less than 0.001% then:

Allowable Blank Residual = Blank Residual

Allowable Blank Residual = 0.007447 mg/ml

If % Blank Residual is greater than or equal to 0.001% then:

Allowable Blank Residual = Density of Acetone x 0.00001 = _____ mg/ml

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